AMENDMENTS TO THE SPECIFICATION

Please amend the specification as described below.

Delete paragraphs 0058-0096.

Replace paragraph 0155 with the following paragraph.

The nucleotide sequences of each of a plurality of GAM oligonucleotides that are described by Fig. 1 and their respective genomic sources and genomic locations are set forth in Tables 1-3, hereby incorporated herein. Specifically, in Table 1, line 778 describes GAM RNA (miRNA) as set forth in SEQ ID NO: 348 is shown as predicted from human.

After paragraph 0155, add the following Table 1, paragraph, Table 2, paragraph, and Table 3.

Table 1

GAM SEQ-ID	GAM NAME	GAM RNA SEQUENCE	GAM ORGANISM	GAM POS
348	GAM353678	CAGCAGCACACTGTGGTTTGTA	Human	A

In Table 2, lines 42112-42207, describes GAM PRECURSOR RNA (hairpin) as set forth in SEQ ID NO: 4233864 and as it relates to Figures 1-8 .

Table 2

GAM NAME	GAM ORGA	PRECUR	PRECURSOR	GAM DESCRIPTION
	NISM	SEQ-ID	SEQUENCE	
GAM	Human	4233	CCTGCTCCCG	Fig. 1 further provides a
353678		864	CCCCAGCAGC	conceptual description of
			ACACTGTGGT	another novel
			TTGTACGGCA	bioinformatically-detected
			CTGTGGCCAC	human oligonucleotide of the
			GTCCAAACCA	present invention referred to
			CACTGTGGTG	here as the Genomic Address
			TTAGAGCGAG	Messenger 353678 (GAM353678)
			GGTGGGGGAGG	oligonucleotide, which
				modulates expression of
				respective target genes whose
				function and utility are known
				in the art. GAM353678 is a
				novel bioinformatically

detectable regulatory, non-

protein-coding, miRNA-like oligonucleotide. The method by which GAM353678 is detected is described with additional reference to Figs. 1-8. The GAM353678 precursor, herein designated GAM PRECURSOR, is encoded by the Human genome. GAM353678 target gene, herein designated GAM TARGET GENE, is a target gene encoded by the target organism as specified in Tables 6-7. The GAM353678 precursor, herein designated GAM PRECURSOR, encodes a GAM353678 precursor RNA, herein designated GAM PRECURSOR RNA. Similar to other miRNA oligonucleotides, the GAM353678 precursor RNA does not encode a protein. GAM353678 precursor RNA folds onto itself, forming GAM353678 folded precursor RNA, herein designated GAM FOLDED PRECURSOR RNA, which has a twodimensional "hairpin" structure. GAM PRECURSOR RNA folds onto itself, forming GAM FOLDED PRECURSOR RNA, which has a two-dimensional "hairpin structure". As is well-known in the art, this "hairpin structure" is typical of RNA encoded by known miRNA precursor oligonucleotides and is due to the full or partial complementarity of the nucleotide sequence of the first half of an miRNA precursor to theRNA that is encoded by a miRNA oligonucleotide to the nucleotide sequence of the second half thereof. A nucleotide sequence that is identical or highly similar to the nucleotide sequence of the GAM353678 precursor RNA is designated SEO ID NO:4233864. and is provided hereinbelow with reference to the sequence listing section. The nucleotide sequence designated SEQ ID NO:4233864 is located from position 7121806 to position 7121896 relative to chromosome 17 on the "-" strand, and overlaps an intergenic region (UCSC.h16.refGene database). Furthermore, the nucleotide sequence designated SEQ ID NO:4233864 is positioned in a region that is conserved

between human, mouse and rat (UCSC.hq16.humorMm3Rn3). A schematic representation of a predicted secondary folding of GAM353678 folded precursor RNA, herein designated GAM FOLDED PRECURSOR RNA is set forth in Table 4 incorporated herein. An enzyme complex designated DICER COMPLEX, an enzyme complex composed of Dicer RNaseIII together with other necessary proteins, cuts the GAM353678 folded precursor RNA vielding a single-stranded ~22 nt-long RNA segment designated GAM353678 RNA, herein designated GAM RNA.Table 5 provides a nucleotide sequence that is highly likely to be identical or extremely similar to the nucleotide sequence of GAM353678 RNA, hereby incorporated herein. GAM353678 target gene, herein designated GAM TARGET GENE, encodes a corresponding messenger RNA, designated GAM353678 target RNA, herein designated GAM TARGET RNA. As is typical of mRNA of a protein-coding gene, GAM353678 target RNA comprises three regions, as is typical of mRNA of a protein-coding gene: a 5' untranslated region, a protein-coding region and a 3' untranslated region, designated 5'UTR, PROTEIN-CODING and 3'UTR, respectively. GAM353678 RNA, herein designated GAM RNA, binds complementarily to one or more target binding sites located in the untranslated regions of GAM353678 target RNA. This complementary binding is due to the partial or full complementarity between the nucleotide sequence of GAM353678 RNA and the nucleotide sequence of each of the target binding sites. As an illustration, Fig. 1 shows three such target binding sites, designated BINDING SITE I, BINDING SITE II and BINDING SITE III, respectively. It is appreciated that the number of target binding sites shown in Fig. 1 is only illustrative and that any suitable number of target binding sites may be present. It is further appreciated that although Fig.

1 shows target binding sites only in the 3'UTR region, these target binding sites may instead be located in the 5'UTR region or in both the 3'UTR and 5'UTR regions. The complementary binding of GAM353678 RNA, herein designated GAM RNA, to target binding sites on GAM353678 target RNA, herein designated GAM TARGET RNA, such as BINDING SITE I, BINDING SITE II and BINDING SITE III, inhibits the translation of GAM353678 target RNA into respective GAM353678 target protein, herein designated GAM TARGET PROTEIN, shown surrounded by a broken line. It is appreciated that the GAM353678 target gene, herein designated GAM TARGET GENE, in fact represents a plurality of GAM353678 target genes. The mRNA of each one of this plurality of GAM353678 target genes comprises one or more target binding sites, each having a nucleotide sequence which is at least partly complementary to GAM353678 RNA, herein designated GAM RNA, and which when bound by GAM353678 RNA causes inhibition of translation of the GAM353678 target mRNA into a corresponding GAM353678 target protein. The mechanism of the translational inhibition that is exerted by GAM353678 RNA, herein designated GAM RNA, on one or more GAM353678 target genes, herein collectively designated GAM TARGET GENE, may be similar or identical to the known mechanism of translational inhibition exerted by known miRNA oligonucleotides. The nucleotide sequence of GAM353678 precursor RNA, herein designated GAM PRECURSOR RNA, its respective genomic source and genomic location and a schematic representation of a predicted secondary folding of GAM353678 folded precursor RNA, herein designated GAM FOLDED PRECURSOR RNA are set forth in Tables 3-4, hereby incorporated herein. The nucleotide sequences of a "diced" GAM353678 RNA, herein

designated GAM RNA, from GAM353678 folded precursor RNA are set forth in Table 5, hereby incorporated herein. The nucleotide sequences of target binding sites, such as BINDING SITE I, BINDING SITE II and BINDING SITE III of Fig. 1. found on GAM353678 target RNA, herein designated GAM TARGET RNA, and a schematic representation of the complementarity of each of these target binding sites to GAM353678 RNA, herein designated GAM RNA, are set forth in Tables 6-7, hereby incorporated herein. It is appreciated that the specific functions and accordingly the utilities of GAM353678 RNA are correlated with and may be deduced from the identity of the GAM353678 target gene inhibited thereby, and whose functions are set forth in Table 8, hereby incorporated herein.

Table 3, lines 1279-1280, shows data relating to the source and location of the GAM oligonucleotide, specifically the GAM PRECRSOR (hairpin) and its position in the genomic sequence of human.

Table 3

GAM NAME	PRECUR SOR SEQ-ID	GAM ORGANISM	SOURCE	STR	SRC-START OFFSET	SRC-END OFFSET
GAM353678	4233864	Human	17		7121806	7121896

Replace paragraph 0156 with the following paragraph.

The nucleotide sequences of GAM PRECURSOR RNAs, and a schematic representation of a predicted secondary folding of GAM FOLDED PRECURSOR RNAs, of each of a plurality of GAM oligonucleotides described by Fig. 1 are set forth in Table 4, hereby incorporated herein. Table 4 lines 2384-2388, shows a schematic representation of the GAM folder precursor as set forth in SEQ ID NO:348, beginning at the 5' end (beginning of upper row) to the 3'

end (beginning of lower row), where the hairpin loop is positioned at the right part of the drawing.

After paragraph 0156, add the following Table 4.

Table 4

GAM NAME	PRE CUR SEQ	GAM ORGA NISM	PRECURSOR SEQUENCE	GAM	FOLDED PRECURSOR	RNA					
	-ID										
GAM 353	423 386	Human	CCTGCTCCCGCCCCAGCAGC	G	C	G	C	T		AC	
678	4		ACACTGTGGTTTGTACGGCA	CCI	CTCCCGCCC	AGCA	CACA	TGTGGTTTG	AC	GGC	T
			CTGTGGCCACGTCCAAACCA	GGA	GGGGGTGGG	TIGI	GTGT	ACACCAAAC	TG	CCG	G
			CACTGTGGTGTTAGAGCGAG GGTGGGGGAGG		AGCGAGA	G	С	C	CA	GT	

Replace paragraph 0157 with the following paragraph.

The nucleotide sequences of "diced" GAM RNAs of each of a plurality of GAM oligonucleotides described by Fig. 1 are set forth in Table 5, hereby incorporated herein. Table 5, line 1276 shows the mature GAM RNA as set forth in SEQ ID NO: 348 as sliced by DICER from the GAM PRECURSOR sequence (hairpin) as set forth in SEO ID NO: 4233864.

After paragraph 0157, add the following Table 5.

GAM NAME	GAM ORGANISM	GAM RNA SEQUENCE	PRECUR SEQ-ID	GAM POS
GAM353678	Human	CAGCAGCACACTGTGGTTTGTA	4233864	A

Replace paragraph 0158 with the following paragraph.

The nucleotide sequences of target binding sites, such as BINDING SITE I, BINDING SITE II and BINDING SITE III that are found on GAM TARGET RNAs of each of a plurality of GAM oligonucleotides that are described by Fig. 1, and a schematic representation of the complementarity of each of these Target binding sites to each of a plurality of GAM RNAs that are described by Fig. 1 are set forth in Tables 6-7, hereby incorporated herein. Table. 6 shows data relating to the SEQ ID NO of the GAM target binding site sequence of the target gene name as bound by the GAM RNA as set forth in SEQ ID NO: 348, Table 6, lines 3688165, 767082, 762322 and 763042 related to target binding site SEQ ID NO: 1810388, 673420, 671402 respectively.

After paragraph 0158, add the following Table 6, paragraph, and Table 7.

Table 6

TARGET BINDING SITE SEO-ID	TARGET ORGANISM	TARGET	TARGET BINDING SITE
0110 000 10	011011112011		SEQUENCE
1810388	Human	MGAT5	CACCATGCTGCTG
673420	Human	SERPINH1	AAACTAGGTGCTGCAG
671402	Human	SERPINH1	ATACCATGATGCTG
671042	Human	SERPINH1	CTATAAAACTAGGTGCTGCAG

Table 7, lines 312839-313773 shows data relating to target genes and binding site of GAM oligonucleotides.

Table 7

GAM NAME	GAM ORGANISM		TARGET BS-SEQ	TARG ET	TARGET REF-ID	TARGET U	TR BINDING SITE DRAW (UPPER:TARGET;LOWER:GAM)	GAM POS
GAM35 3678	Hum	CAGCAGCA CACTGTGG TTTGTA	AAACCAAA CTTATGCA GCTG	nup C	NC_004431 f rom 27953 90 to 27966 31 (+)	Escher 3 ichia coli CFT073	A C TA A AAACCA A T TGC GCTG TTTGGT T A ACG CGAC ATG G C C- A	A
GAM35 3678	Hum	CAGCAGCA CACTGTGG TTTGTA	AAACCAAA CTTATGCA GCTG	nup C	NC_004741 f rom 24940 19 to 24952 21 (+)	Shigell a fle 3 xneri 2a str . 2457T	A C TA A AAACCA A T TGC GCTG TTTGGT T A ACG CGAC ATG G C C- A	Α
GAM35 3678	Hum	CAGCAGCA CACTGTGG TTTGTA	AAACCCTG CTGCG	rel A	NC_000962 f rom 29078 24 to 29101 96 (-)	Mycobac 3 teriu m tubercu los is H37Rv		A

GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	AAACCCTG CTGCG	rel NC_002945 f ter rom 28752 sul 74 to 28776 box 46 (-) AF; 22/	cobac fiu m 3 C C A fis AAACC TGCTGC G A fis TITIGG ACGACG C fis ATG TGTCAC A fig TGTCAC A
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	AAACCCTT TCTGCTGC TT	yab NC_004431 f Esc 0 rom 614 col 89 to 62148 CF1	197 cheri 3 C TIC- T A cla AAACC T TGCTGCT A cli TTTGG G ACGACGA coro T TCAC C
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	AAACCGAT GCAGTGCG GCTG	amt NC_004337 f Shi B rom 4080 kne 59 to 40934 2a 5 (+) . 3	gell 3 AT CAG G tle AAACCG G TGC GCTG A eri TITGGT C ACG CGAC str ATG GT AC- A
GAM35 Hum CAGCAGCA 3678 an CACIGIGG		amt NC_004741 f a d rom 4078 xne 60 to 40914 2a	igell lle 3 AT CAG G ri AAACCG G TGC GCTG A str TITIGGT C ACG CGAC 457T ATG GT AC- A
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	AAACCGAT GCCGTGCG GCTG	amt NC_004431 f Esc b 16 to 54890 cF1 2 (+)	cheri 3 AT CCG G La AAACCG G TGC GCTG A Li TITGGT C ACG CGAC 1073 ATG GT AC- A
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	AAACCGCC CCCAGTCT GCTG	dsd NC_003197 f Sal A rom 40044 yph 53 to 40057 ium 75 (+) T2	Limone 3 CCCCAG - A t AAACCGC T A AAACCGC T A AAACCGC ATG TTGGTG h L A GACGAC ATG TCAC C
3678 an CACTGTGG	CTTGCCGC	gad rom 48716 put 25 to 48729 put	AAACCG T TGC GCTG
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	AAACGAAT TGAATCAT GCCGCTG	aro NC_003116 f Nei A 02 to 15588 dis 03 (+) 249	ATG GTC C A me AAAC A TGC A mgti GCTG TTTG T 2 AGG CGAC ATG ll G GTCAC A resini 3 G AAAC ACG GCTGCTG TTTG CACA cose GCTGCTG TTTG AGG GCTGCTG TTTG AGG GCTGCTG TTTG cose AAC ACG cose AGG GCTGCTG TTTG
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	AAACGACG GGCTGCTG	ruv NC_003143 f Yer B rom 23364 a p 49 to 23374 a p 53 (+)	rsini 3 GG AAAC A es ngt cgacgac atg G caca
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	AAACGACG GGCIGCIG	31 to 24830 tis	KIM TGT CGACGAC ATG G
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	TCCTGCTG	yei NC_004431 f Esc E rom 15586 col 41 to 15591 cF1	cheri 3 G C ta AAAC ATA T CTGCTG A ti TITG TGT A GACGAC TO73 ATG G CAC C
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	AAACGCAT GTTCATGC GCTG	van NC_002516 f Pse D rom 55041 aer 20 to 55050 aer 73 (+) Add	eudom 3 G T CA - ss 3 AAAC CATG T TGC A rugin GCTG TTTG GTGT A a P ACG CGAC ATG - C L C- A
GAM35 Hum CAGCAGCA 3678 an CACTGTGG	AAACGCTC CGTATACT	fts NC_000922 f Chl Y rom 11154 oph	Lamyd 3 GC CCGTATAC A

	GCTGCTA			TGCTGCT TTTG G ACGACGA ATG GT TCAC C
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	AAACGCTC CGTATACT GCTGCTA	fts NC_002491 f rom 11131 27 to 11139 99 (-)	Chlamyd 3 ophil a pneumon iae J138	GC CCGTATAC A AAAC T A TGCTGCT TITG G ACGACGA ATG GT TCAC C ATCGACAGT
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	AAACTAAT CGACAGTT GCTGCG	rbs NC_004337 f R 08 to 39487 00 (+)	a fle xneri 2a str	- AAACTA A TGCTGC G A TTTGGT ACGACG C ATG
GAM35 Hum CAGCAGCA 3678 an CACTGTGG	AAACTAAT CGACAGTT GCTGCG	rbs NC_004741 f R rom 38245 94 to 38255 77 (-)	a fle 3 xneri 2a str . 2457T	ATCSACAGT - AAACTA A TGCTGC G TTTGGT ACGACG C ATG
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	AAACTAGG TGCTGCAG	SER PIN NM_001235 H 1	3 Human	AAACTA G TGCTGC G A TTTGGT T ACGACG C
GAM35 Hum CAGCAGCA 3678 an CACIGIGG TIIGIA	AAACTCAG GCTGGCAA GCTGCTG	aro rom 15575 H 27 to 15585 73 (-)	a fle xneri 2a str . 301	AAAC CA G IG A GCIGCIG TITG GI I AC CGACGAC AIG - G CA
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TITGTA	AAAGCTGC TGCTT	zra NC_003197 f rom 43877 27 to 43881	Salmone 3 lla t yphimur	G T AAA C TGCTGCT A
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TITGTA	AAATCAGT TGTACTTG TTGCTG	cys NC_003197 f m 25516 51 to 25525 62 (-)	Salmone 3 11a t yphimur ium L T2	ATG G TGTCAC C GTIGTACT T AAAATCA TG A TGCTG TTTGGT AC ACGACC ATG GTCAC G AT TGCTGCTG TT G ACGACGAC ATG T G TCAC C ACGACGAC ATG T G TCAC
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TITGTA	AACATTGC TGCTG	rbs NC_004431 f rom 44392 60 to 44402 52 (+)	Escheri 3 chia coli CFT073	AT TGCTGCTG TT G ATG TCAC
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	AACTGCTG CTC	oxy NC_003197 f rom 43430 80 to 43439 97 (+)	Salmone 3 lla t yphimur ium L T2	AA C TGCTGCT A TT G ACGACGA ATG T GTGTCAC C
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	AACTGCTG CTC	oxy NC_003198 f rom 36072 R 04 to 36081 21 (-)	Salmone lla e nterica 3 ente rica serovar Typhi	AA C TGCTGCT A TT G ACGACGA ATG T GTGTCAC C
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	AACTGCTG CTC	oxy NC_004631 f rom 35928 R 64 to 35937 81 (-)	Salmone lla e 3 nterica ente rica serovar	AA C TGCTGCT A TT G ACGACGA ATG T GTGTCAC C

		Typhi Ty2	
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	AAGCCGGT TGCGGTGC TGCTG	NC_003116 f Neisser AAGCCG AAGCCG TGCTGCTG TGCTGCTG TGCTGCTG TGCTGCTG TTGCT TTGCT ACGACGAC ATG GTACK	¥
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TITGTA	AATCCACT CCGTGTTG CTG	Nc_003198 f nertca Same	¥
GAM35 Hum CAGCAGCA 3678 an CACTGIGG TTIGTA	AATCCACT CCGTGTTG CTG	Salmone 11a e g1_ NC_004651 fn terica 3 T TCC T AA F rom 41292 ente CCAC GTG TGCTG TT J 15 to 41316 rica GSTG CAC ACGAC ATG T 62 (+) SECOVAT TCA G CAC ACGAC ATG T TyPh Ty2	
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA		NC_004431 f Becheri 3	¥
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	ACATGCTG CTT	Ohigell NC_004741 f a fle 3 TGCTGCT NUP FOR 24940 kmeri A C A TGCTGCT 19 to 24952 2a str T G T ACGACGA 21 (+) . 24571 ATG IT G GTCAC C	¥
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	GTACTGCT	ho NC_000962 f Mycobac 3 G TAC	¥
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	ACGATGGT GTACTGCT GCTT	Mycobac Myco	A.
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA		Salmone Salmone Salmone	¥
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	ACTGCTGC TC	Salmone Lia e C P C C P C C P C C	¥
		rec NC_002677 f Mycobac 3 T TTAG 3 G rom 20147 terium AGA ATG TG	¥

TTTGTA		54 (-)		TGCTGCTG TTT TGT AC ACGACGAC ATG GG C
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	AGACCATG AAGAACTG A	rel NC_000962 f 1 A rom 29078 24 to 29101 96 (-)	Mycobac 3 teriu m tubercu los is H37Rv	AA AACTG AGACCATG G A GCTGCTG TTTGGTGT C CGACCAC ATG CAA
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	AGACCATG 1 AAGAACTG 2 GCTGCTG	rel NC_002945 f rom 28752 74 to 28776 46 (-)	Mycobac teriu m 3 bovis subs p bovis AF21 22/97	AGACCATG G AGCTGCTG TITGGTGT C CGACGAC ATG CAA
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	AGCITGCI GCIG	c 50 to 42908	a pes tis	TGCTGCTG TT GG " ACGACGAC ATG T TGTCAC
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	AGCTTGCT G	glp NC_004088 f 7 C 77 to 45604 7 (+)	Yersini 3 a pes tis KIM	TGCTGCTG TT GG A ACGACGAC ATG T
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	ATACCAAG GCTGCTG	fts NC_000922 f rom 11154 1 27 to 11162 99 (-)	Chlamyd ophil a 3 pneumon iae CWL029	ACCA GCTGCTG T A TGGT CGACGAC ATG T
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	ATACCAAG GCTGCTG	fts NC_002491 f Y rom 11131 Y 27 to 11139 1 99 (-)	Chlamyd 3 ophil a pneumon iae J138	ACCA GCTGCTG T A TGGT CGACGAC ATG T
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	ATACCATG E	SER PIN NM_001235 : H 1	3 Human	T A A A A A A A A A GCA TG TGCTG T TGGT AC ACGAC ATG T GTCAC G
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTIGTA	ATTICIGC I	aro NC_004342 f : D rom 481 : D 28 to 48832 (-)	Leptosp ira i nterrog 3 ans s erovar lai s tr. 56601	A TC TGCTGCT A T GG ACGACGA ATG IT TGTCAC C
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	0.0	7 (+)	e	A- C C CAAA ACGGT TGC GCTG GTTT TGTCA ACG CGAC AT GG C A
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CAAAAGCT : TTTCGCTT : GCGGCTG	zra NC_003197 f rom 43877 27 to 43881 82 (-)	Salmone 3 lla t yphimur ium L	GCT TTCGCT G CAAA A T TGC A GCTG GTTT T A ACG CGAC AT GG
	CGGCIG	glp NC_004310 f D 63 to 21227 4 (+)	Brucell 3 a sui s 1330	GTCGTCCTG G CAAA A TGC A GCTG GTTT T ACG CGAC AT GG GTCAC A
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CAAACAGG & CIGCIGCI G A	avt NC_004337 f : A rom 37211 : 75 to 37225 :	Shigell 3 a fle xneri	G C CAAAC A A G TGCTGCTG GTTTG T T ACGACGAC AT G

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33 (+) 2a str G CAC
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                                                                                                                                                                                                                                                                                     Shigell
GAM35 Bum CAGCAGCA CAAACAGG avt NC_004741 f a fle 3 -- - G C-- CAAAC A 3678 an CACCAGGA CTGCIGCI G A 70m 40526 xneri A G TGCTGCTG GTTTG 25 to 40539 2a str T I ACGAGCAC AT G
                                                         TTTGTA
                                                                                                                                                                                                           38 (-) . 2457T G CAC
 GAM35 Hum CAGCAGCA CARACCAG sel NC_002947 f Peeudom 3 -- GC G-- GC G-- GAACCAG CAGCAGCA G T.CTGCTG A STORM SE21 onas CARACCA G T.CTGCTG A TITTGTA TG S (5 (+) CARACCA G T.CTGCTG A GTTTGGT C A GACGAC G T.CTGCTG A GACGAC G T.
GAM35 Hum CAGCAGCA 3678 an CACTGTGG CAMACCAT TOTTGTA CATGCTG A 7 to 90029 achomat GTTGTA AC ACCAC 5 (+) is AT GTCAC G
GAM35 Hum CAGCASCA CAAACCCC dna ^{\circ} AGACCCC ASTRONOM ACTION A
                                                                                                                                                                                                                                                                                     Leptosp
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GAM35 Hum CAGCASCA CAAACIC are NC_004342 fintering 3 -- C TI IC T
3678 an CACTGIGG THICTIC D to nom 481 and 5 CAAACI T I I CIGGIG A
TITISTA IGCIG (-) Laid S AT I IC C- C
                                                                                                                                                                                                                                                                                       tr.
                                                                                                                                                                                                                                                                                       Salmone
60 (+) rica TCAC
GAM35 Hum CAGCAGCA CAAAGCCG amt NC_004431 f Escheri 3 -- G CGC -- G GAAGCCG amt rom 5476 chia CAAA G TGC GCTG A 3678 am CACTGTGG CGTGCGCT B 16 to 54890 coli GTTT G C ACG CGAC 2 (+) CFT073 AT G TGT AC A
                                                                                                                                                                                                                                                                                    Shigell
 GAM35 Hum CAGCAGCA CAAAGCCG amt NC_004741 f a file 3 - G CGC - G TGC GCTG A 1678 an CACTGTGG CTG G F Cot 4078 xneri CAAA C G TGC GCTG A 171G GCTG A 17
                                                                                                                                                                                                           6 (+) . 2457T AT G TGT AC A
 GAM35 Hum CAGCAGCA CAAAGCCT pil NC 002947 f Pseudom 3 -- G C TT TTCGG A
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	TTTTTCGG GCTGCTG	34 to 58179 putida GCTGCTG GTTT G G 44 (-) KT244 0 A CGACGAC AT G
	CATGTAAT GCTTGCTG	62 (-) T2 TC
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CAAATCCC CAGTIGIG CIG	NC_004431 f Echeri 3 C CAGT Step
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	TACTGCTT	H rom 15575 xneri CTG GTT GTG 27 to 15585 2a str ACGA GAC AT TG 73 (-) .301 TCAC
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CAACACTT TGCGCTG	NC_003197 f Salmone 3 TIT- G GCTG A 1942 lia t CAA C AC TGC GCTG A 0 1 to 19499 ybihumr GTT G TG ACG CGAC 5 (-) TZ AT T G TCAC A
GAM35 Hum CAGCAGCA 3678 an CACIGIGG TITGIA	CAACACTT TGCGCTG	Salmone Salm
		Salmone
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CAACAGCA GTTGCTGC TG	ace NC_002947 f Fseudom 3 GCAGI CAA C rom 51847 onas a TGCTGCTG GTI G A 42 to 51864 put.lda a TGCTGCTG GTI G F57 (-) KT244 0 GTCAC
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CAACAGIT TCTTGGTC TGCTG	fmt NC_002745 f 00000 3 GTI CTIGG - CAA C A T T T A CTIGG G ST G T G T G T G T G T G T G T G T
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CAACAGTT TCTTGGTC TGCTG	$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CAACAGTT TCTTGGTC TGCTG	$\begin{array}{cccccccccccccccccccccccccccccccccccc$

		В	aureus MW2	
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CAGCACTG			C CAGCAC - CAA CCAC TGCTGC A G GTT GGTG ACGACG C AT T TCAC A
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CAACCGCT GCTGCTG	pho NC_000962 Y2 rom 913 56 to 9141 7 (-)	H37Rv	CGC TGCTGCTG GTT A GGTG ACGACGAC AT T TCAC
GAM35 Hum CAGCAGCA 3678 an CACIGIGG TIIGTA	GCIGCIG	9 (-)	AF21 22/97	CCGC TGCTGCTG GTT A GGTG ACGACGAC AT T TCAC
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CAACCGGT GCTGCG	dad NC_004431 X 06 to 1477 76 (+)	f Escheri 3 chia coli CFT073	CAA CCG TGCTGC G GTT GGT ACGACG C AT T GTCAC A
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	GGTGATGC	fha NC_002929 L rom 3085 L 65 to 3098 55 (+)	8 IIA P 14 ertussi	C GTGA - CAA CCGT G TGCT A CTG GTT GGTG C ACGA GAC AT T T AC C
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CAACCTGC GCTG	spe NC_003197 D rom 1949 5 (-)	f Salmone 3 12 lla t 2 yphimur 1 ium L T2	CAA CC IGC GCIG A GIT GG ACG CGAC AI I TIGICAC A
GAM35 Hum CAGCAGCA 3678 an CACIGIGG TITGTA	CAACCIGC GCTG	spe NC_003198 rom 190 89 to 1971 3 (-)	Typhi	CAA CC TGC GCTG A GTT GG ACG CGAC AT T TGTCAC A
GAM35 Hum CAGCAGCA 3678 an CACIGIGG TITGIA	CAACCIGC GCIG	spe NC_004631 prom 196 80 to 1971 4 (-)	Salmone 11a e f nterica 3 3 ente 7 rica serovar Typhi Ty2	CAA CC TGC GCTG A GTT GG ACG CGAC AT T TGTCAC A
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA		21 (+)	s	CGA CAA CCA GCTGCTG GTT A GGT CGACGAC AT T- GTCACA
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CAAGCCAA TCTGCTG	sse NC_004431 B rom 2922 56 to 2923 41 (-)	2 coli	CAAGCCA T CTGCTG A GTTTGGT A GACGAC
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CAAGCCCG CTTGCTGT G	aer NC_002947 rom 2406 -2 96 to 2408 61 (-)	15 PULLUA	CGCT A CAAGCC TGCTG TG GTTTGG ACGAC AC AT TGTCAC G

GAM35 Hum CAGCAGCA 3678 an CACIGIGG TITGIA	CAAGCCTG CGCTG	lpp NC_000962 f rom 22912 67 to 22919 23 (+)	Mycobac 3 teriu m 3 tubercu los is H37Rv	CAAGCC TGC GCTG A GTTTGG ACG CGAC AT TGTCAC A
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA		lpp NC_002945 f rom 22751 82 to 22758 38 (+)	22/97	CAAGCC IGC GCIG A GTITGG ACG CGAC AT IGTCAC A
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CAAGCTGC TGCTG	ris NC_002929 f rom 37652 57 to 37659 91 (-)	Bordete 3 lla p ertussi s	CAAGC A TGCTGCTG GTTTG A ACGACCAC AT GTGTCAC GG
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CAAGGACC ATGCGCTG	acc NC_002929 f rom 9264 C 07 to 92777	lla p ertussi	CAA ACCA TGC A GCTG GTT TGGT ACG CGAC AT
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CAAGGCAA AGGTCTGC TG	rps NC_002947 f rom 7070 68 to 70734 6 (-)	Pseudom 3 onas putida KT244 0	GTCAC A G A G CAAG A CA AG T CTGCTG GTTT A GT TC A GACGAC AT G G AC C
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CAATAACA ATGCAGCT G	fmt NC_002745 f B (m NC_02745 f 3 r rom 22181 r 45 to 22255 p) 90 (-)	Staphyl ococc 3 us 3 aureus su bsp. aureus N315	T A A A CAA A CA TGC GCTG A GTT T GT ACG CGAC AT G GTCAC A
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CAATAACA ATGCAGCT G	fmt NC_002758 f B(m NC_02758 f 3 r rom 22879 3 t 22953 p) 80 (-)	Staphyl occocc 3 us aureus su bsp. aureus Mu50	T A A A CAA A CA TGC GCTG A GTT T GT ACG CGAC AT G GTCAC A
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CAATAACA ATGCAGCT G	tru nca NC_003923 f t rom 22380 s ed 83 to 22401 fmt 43 (-) B	us 3 aureus su bsp. aureus	T A A A CAA A CA TGC GCTG A GTT T GT ACG CGAC AT G GTCAC A
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CAATAGCG CCTGCTGC TG	nup NC_004337 f c NC_004337 f rom 25158 42 to 25170 83 (+)	Shigell 3 a fle xneri 2a str . 301	T GC CC CAA A G IGCIGCIG GTI A T C ACGACGAC AT IGG GT AC
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CAATAGCG CCTGCTGC TG	nup NC_004431 f rom 27953 c 90 to 27966 31 (+)	Escheri 3 chia coli CFT073	T GC CC CAA A G TGCTGCTG GTT A T C ACGACGAC AT TGG GT AC
GAM35 Hum CAGCAGCA 3678 an CACTGTGG	CAATAGCG	NC 004741 f	Shigell a fle 3	T GC CC CAA A G TGCTGCTG GTT T C ACGACGAC AT

	CAATATAG AAGCTGCT GCTA	NC_000922 f op. def rom 12217 pn 35 to 12222 ia 95 (+) C	WL029 AT TG G CAC C
GAM35 Hum CAGCAGCA 3678 an CACIGIGG TIIGIA	CAATATAG AAGCTGCT GCTA	29 (+) ja	138 AT TG G CAC C
	CAATCACC GGGCCGAT GCGGCTG	glc NC_004431 f Es c nom 35428 ch. 71 to 35436 cp 95 (+)	cheri 3 T - C G CCGA G ia CAA C AC G G TGC A ii GCTG GTT G TG C C T073 ACG CGAC AT T G T A A
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CAATCAGG GATACTGC TG	pta rom 8916 on 25 to 89371 pu 2 (-)	cida GITC A GACGAC AT I 2440 GG AC C
GAM35 Hum CAGCAGCA 3678 an CACIGIGG TIIGIA	CAATCCCC GCTTCCTG CTG	A 36 to 12663 er 79 (+) s	TCCC
	CAATCCCG GCCATTTG CTCTG	ruv NC_003143 f Yes B 23364 a 1 49 to 23374 a 1 53 (+)	CAC C
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TITGTA	CAATCCCG GCCATTTG CTCTG	Tuv NC_004088 f Yes B rom 24820 a g 31 to 24830 til 35 (-)	S KIM ACGA GAC AT T TG CAC C
	CAATCGCA GCACTGGT GCTG	C 42 to 25170 xn 83 (+) 2a	str GTT G GT AC ACGAC 301 AT T - GTCAC G
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TITGTA	CAATCTCA CTTTCTGC GCTG	rps NC_002947 f on. T fom 7070 pu 68 to 70734 pt 6 (-)	
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CACACAGG TCATCCTT GCGCTG	prc NC_002677 f My. A rom 15765 te: 53 to 15773 le: 50 (+)	prae ACG CGAC AT T G G CAC A
GAM35 Hum CAGCAGCA 3678 an CACIGIGG TIIGIA	CACATGTT GTACATGC TGCTT	C from 25158 xn 42 to 25170 xn 83 (+) 2a	eri TGCTGCT GT G TGT str AC ACGACGA AT TT G
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TITGTA	CACCAATG CTCCTG	23 (1) 10	oobad 3 C crium 3 A C crium CA CCA TGCT CTG bercu GT GGT ACGA GAC s is AT TT GTCAC C
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CACCAATG CTCCTG	lpp NC_002945 f My I rom 22751 te 82 to 22758 bo	cobac 3 A C A riu m CA CCA TGCT CTG vis GT GGT ACGA GAC

		38 (+)	subs p bovis AF21 22/97	AT TT GTCAC C
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CACCACCG om CTAACTGC G TGCG	NC_004431 f rom 16245 77 to 16255 33 (+)	Escheri 3 chia coli CFT073	C CTAAC - CA CCAC G A TGCTGC G GT GGTG C ACGACG C AT TT T AC A
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CACCACCT Ph CCTGCTG V	n NC_003198 f rom 4715 75 to 47236 6 (-)	serovar Typhi	C C CA A CCAC I CIGCIG GT A GGIG A GACGAC AT TI TCAC C
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CACCACCT Ph CCTGCTG V	n NC_004631 f rom 25087 35 to 25095 26 (+)	Salmone lla e nterica 3 ente rica serovar Typhi Ty2	
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CACCACGT pi AGTGCTTC T TG	NC_002947 f rom 58169 34 to 58179 44 (-)	Pseudom 3 onas putida KT244 0	CA CCACG GIGCI CIG GI GGIGT CACGA GAC
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TITGIA	CACCAGCG OX CCTGCGCT G R	y NC_003197 f rom 43430 80 to 43439 97 (+)	Salmone 3 11a t yphimur ium L T2	GC CC - CA A CCA G IGC GCIG GT GGT C ACG CGAC AT TT GT AC A
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CACCAGCG ox CCTGCGCT G R	y NC_003198 f rom 36072 04 to 36081 21 (-)	Salmone lla e 3 nterica ente rica serovar Typhi	GC CC - CA A CCA G TGC GCTG GT GGT C ACG CGAC AT TT GT AC A
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CACCAGOG ox COTGOGOT G R	04 10 33537	Salmone 11a e nterica 3 ente rica serovar Typhi Ty2	GC CC - CA A CCA G TGC GCTG GT GGT CC AGG GGAC AT TT GT AC A
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CACCATGC MG TGCTG T5	A NM_002410	3 Human	CCA TGCTGCTG GT A GGT ACGACGAC AT TT GTCAC
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CACCATTG ip	a NC_004741 f a rom 20232 05 to 20248 48 (+)	Shigell a fle 3 xneri 2a str . 2457T	CA CCAT TGCTGC G A GT GGTG ACGACG C AT TT TCAC A
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TITGTA	CACCATTG H9 CTGCCG 8	a NC_004337 f . rom 14220 64 to 14237 79 (-)	a fle xneri	CA CCAT TGCTGC G A GT GGTG ACGACG C AT TT TCAC A

			. 301	
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CACCATTG 5.	it NC_004337 f rom 14053 60 to 14062 17 (-)	Shigell 3 a fle xneri 2a str . 301	CA CCAT TGCTGC G A GT GGTG ACGACG C AT TT TCAC A
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA		it NC_004741 f rom 19046 66 to 19055 23 (+)	2a str	CA CCAT TGCTGC G A GT GGTG ACGACG C AT IT TCAC A
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CACCATIT di CTGCCGCT G E	47 (+)		TTC- C CA A CCAT TGC GCTG GT GGTG ACG CGAC AT TT TCAC A
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CIGCCGCI G A	28 (-)		TIC- C CA A CCAT TGC GCTG GT GGTG ACG CGAC AT TT TCAC A
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CIGCCGCI G A			GGTG ACG CGAC AT TT TCAC A
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA		78 (-)		TTC- C CA A CCAT TGC GCTG GT GGTG ACG CGAC AT TT TCAC A
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA		76 (+)	022010	
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CACCCAGG n: TGTCGCTG T CTG	ic NC_000962 f rom 31666 81 to 31677 99 (+)	Mycobac 3 teriu m 3 tubercu los is H37Rv	C- G TGTC CA A CCA G GCTGCTG GT A GGT T CGACGAC AT TT G CACA
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA			22/97	C- G TGTC CA CCA G GCTGCTG GT A GGT T CGACGAC AT TT G CACA
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	TGCTGCTC A	57 to 37659 91 (-)	ertussi s	T TC C A CA CC T TGCTGCT GT GG G ACGACGA AT TT T TCAC C
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA				C TGTGTT - CA GCCATA TG A TGCTG GT TGGTGT AC ACGAC AT T CAC G
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CACGCCAT HEATGTGTTT B	pa NC_004337 f 9. rom 14220 64 to 14237 79 (-)	Shigell 3 a fle xneri 2a str . 301	C TGTGTT - CA GCCATA TG A TGCTG GT TGGTGT AC ACGAC AT T CAC G
GAM35 Hum CAGCAGCA 3678 an CACTGTGG	CACGCCAT S:	it NC_004337 f	Shigell 3	C TGTGTT - A CA GCCATA TG

TTTGTA				TGCTG GT TGGTGT AC ACGAC AT T CAC G
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA				C TGTGTT - CA GCCATA TG A TGCTG GT TGGTGT AC ACGAC AT T CAC G
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA		00 (+)	301	A T TACC CAGA CAC GT A GCTGCTG GTTT GTG CA CGACGAC AT G T CA
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA				A T TACC CAGA CAC GT A GCTGCTG GTTT GTG CA CGACGAC AT G T CA
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA		,		A T TACC CAGA CAC GT A GCTGCTG GTTT GTG CA CGACGAC AT G T CA
GAM35 Hum CAGCAGCA 3678 an CACIGIGG TIIGIA			. 3UL	G G AACAT CA AA CATG TGCTGCTG A GT TT GTGT ACGACGAC AT - G CAC
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA		NC_004741 f rom 38245 94 to 38255 77 (-)	a fle 3 xneri 2a str . 2457T	G G AACAT CA AA CATG TGCTGCTG A GT TT GTGT ACGACGAC AT - G
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CAGACGAT fer CTCCTGCT G C	NC_002516 f rom 46535 08 to 46543 05 (-)	Pseudom 3 onas aerugin osa P A01	G C C CAGAC AT T CTGCTG A GTTTG TG A GACGAC AT G TCAC C
GAM35 Hum CAGCAGCA 3678 an CACTGTGG	CAGACTCA pil GCTGCTGC T TC	NC_002947 f rom 58169 34 to 58179 44 (-)	Pseudom 3 onas	I GC C A CAGAC CA IGCIGCI GITIG GI ACGACGA AI - GICAC C
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CAGCAGGC eII	NC_003198 f rom 45373 12 to 45375 33 (+)	Salmone lla e 3 nterica ente rica serovar Typhi	G C T CAG C A G T TGCTGCTG GTT G T T A ACGACGAC AT T G G C C
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	TTTGCTGC CIT	from 45201 21 to 45203 42 (+)	ente rica serovar Typhi Ty2	
GAM35 Hum CAGCAGCA 3678 an CACTGTGG	CAGCCACA gad GCTGCTG	NC_002947 f rom 48716	Pseudom 3 onas	A CAG A CCAC GCTGCTG GTT

TTTGTA		25 to 48729 put 29 (+) KT2	ida GGTG CGACGAC AT T
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CAGCCAGG TTTTGCTC TG	phn NC_003198 f nte V rom 4715 ent 75 to 47236 ric 6 (-) ser Ty	mone e g - G I I - rica CAG CAG I TGCI CIG e GIT GOI T A ACGA GAC caa GIT GOI T A ACGA GAC covar A I G C C C phl mone
		phn NC_004631 fn rom 25087 ent V 35 to 25095 ric 26 (+) ser Ty	ne urica 3 G T T A ce CAG CCA G T TGCT CTG A ca GTT GGT T A ACGA GAC ovar AT G C C C phi
		75 (17)	dete 3 G G CCTCG CAG A D GTT G T T CUSSI ACGACGAC AT T G G CAC
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CAGCGCTT GGTGATGC TGCTG	uhp NC_003143 f Yer A 90 to 45233 a p 80 (-)	rsini 3 CG T A CAG C TGGTG TGCTGCTG GTT A G GTCAC ACGACGAC AT TG T -
GAM35 Hum CAGCAGCA 3678 an CACIGIGG	CAGGCGCA	cys NC_002947 f Pse rom 3151 ona Q 84 to 31598 put 4 (+)	rudom 3 G G CAGGC CA A 1.1 G GGTGTGCTGCTG GTTTG GT A 1.1 CACACGACGAC AT - G G CAGGC CA A 1.1 CACACGACGAC AT -
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CATACCTC CCGCACTG CTGCCG	pbp NC_002947 f Pse prom 43237 ona G 07 to 43246 put 33 (+) KT2	eudom 3 T T CCGCAC C is CA ACC C TGCIGC A ida G GT IGG G 444 O ACGACG C AT T T TCAC A
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CATATOTS CTGCTG	NC_000907 f Hae ung rom 186 ilu 76 to 19335 inf (+) zae	moph 3 T CA ATC A Studen ACGACGAC AT T TGTCAC
GAM35 Hum CAGCAGCA 3678 an CACTGIGG TITGIA	CATCCACA CGCTGCTG	2 (17	sudom 3 TC CA A s
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CATCCATA TCGCCATT GCTGGTG		theri 3 T- TC CCAT a. G CA CCATA G A i. TGCTG TG GT GGTGT 1073 C ACGAC AC AT TT CA G
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA		glp NC_003143 f Yer C rom 42896 a p 50 to 42908 tis	- TCG G CGCGC - Sini
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CATCGCGG GCGCGCTG CTGCTC	glp NC_004088 f Yer C 77 to 454604 tis 7 (+)	rsini 3 TCG G CGCGC C CA CG G A CES TGCTGCT GT GT T ACGACGA AT TTG G
			CAC C

	GCTGCTT	14 to 14265 leprae
TTTGTA	GCTGCTT	pon NC_002677 f Mycobac 3 T GT GTGGA A
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TITGTA	CATGICGG TGGIGGGT GCTGCTT	NC_002677 f Mycobac 3
GAM35 Hum CAGCAGCA 3678 an CACIGIGG TITGIA	CATGICGG IGGIGGGI GCIGCII	78 (-) C ACGACGA AT T GT AC C
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TITGTA		omp rom 16245 chia c ccG TGCTGGT A G 77 to 16255 chia G GST ACGACGA 33 (+) CFT073 AT TIT GTCAC C
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CCCTCGGT GCTGCTG	fih NC_002929 f Bordete 3 T C CC A B rom 14417 lla p CGG TGCTGCTG G GG A 67 to 14429 ertussi GTC ACGACGAC AT TTT T 21 (+) S AC
	CGATGCTG TG	-2 96 to 24085 putida GTTTGGTG TG ACGAC AC 61 (-) KT244 0 AT T AC G
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TITGTA	CGAACGTG CTGCTG	Pro NC_002677 f Mycobac 3 G CGAAC AT A 53 to 15773 leprae A GGAGGAC AT GGGGGAC AT GGGGGAC AT GGGGGAC AT GGGGAC AT GGGAC AT GGGAC AT GGGAC AT GGGAC AT GGAC AT GGAC AT GGGAC AT GGAC AT
GAM35 Hum CAGCAGCA 3678 an CACIGIGG TIIGIA	GCGTGGTG	B 20 to 55050 osa P T A G
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TITGTA	CGACTGCT GCTG	Poh NC_002516 f Pseudom 3 CGA C A 7451 onsa TGCTGCTG GTT G A 20 to 47455 serugin ACGACGAC AT T 50 CGA C A 750 CGA C A
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TITGTA	CGAGCGAT GCTGCTT	dsd NC_003197 f Salmone 3
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	GTAGTGCT	
36/6 an CACIGIGG	CGAGGGAT GTAGTGCT GCTC	NC_002945 fterlum 3

22/97

GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CGCCAGAT TGCTGCTG	yab O	NC_004431 f rom 614 89 to 62148 (-)	Escheri 3 chia coli CFT073	CCA A TGCTGCTG GT A GGT T ACGACGAC AT TT G CAC
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CGCCGGTT GCTGCTG	pch A	NC_002516 f rom 47451 20 to 47465 50 (+)	Pseudom 3 onas aerugin osa P A01	CCG TGCTGCTG GT A GGT ACGACGAC AT TT GTCAC
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CGCCGGTT GCTGCTT	fep C	NC_002516 f rom 46535 08 to 46543 05 (-)	Pseudom 3 onas aerugin osa P A01	CG CCG TGCTGCT A GT GGT ACGACGA AT TT GTCAC C
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CGCTGCTG CTG	liv H	rom 11447 29 to 11456	Bordete 3 lla p ertussi	TGCTGCTG GT G A
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CGGAATTT GCTGCTG	yci E	NC_004431 f rom 15586 41 to 15591 47 (-)	Escheri 3 chia coli CFT073	T CGGA AT TGCTGCTG GTTT A TG ACGACGAC AT GG TCAC
			2.2		T- CG GTGG C CG CCACG G A TGCTGCT GT GGTGT C ACGACGA AT TT CA C
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CGTCCATC TTGCTGCT C	pbp G	NC_002947 f rom 43237 07 to 43246 33 (+)	Pseudom 3 onas putida KT244 0	CG CCAT TGCTGCT A GT GGTG ACGACGA AT TT TCAC C
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CGTCCTGC TGCTG	aco R	NC_002516 f rom 46395 01 to 46413 78 (-)	Pseudom 3 onas aerugin osa P A01	TE TE TO THE TOTAL TO THE TOTAL TOTA
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CGTGCCTG CTGCTC	ace K	NC_002947 f rom 51847 42 to 51864 57 (-)	Pseudom 3 onas putida KT244 0	T C A CG GCC TGCTGCT GT TGG ACGACGA AT T TGTCAC C
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CTAAAGTG CTGCTG	rec G	NC_002677 f rom 20147 23 to 20169 54 (-)	Mycobac 3 teriu m leprae	T G C AA A TGCTGCTG G TT T ACGACGAC AT T GG GTCAC
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CTAACACG CGCTGCTC TG	ung	NC_000907 f rom 186 76 to 19335 (+)	Haemoph 3 ilus influen zae R d	T - CGC - C AAC ACG TGCT CTG G TTG TGT ACGA GAC AT T G CAC C
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CTAACCTG TATCTGCT G	cII	NC_003198 f rom 45373 12 to 45375 33 (+)	Salmone lla e nterica ente rica serovar Typhi	I IGTA C AACC T CIGCIG G A TIGG A GACGAC AT I TGTCAC C
GAM35 Hum CAGCAGCA	CTAACCTG	cII	NC_004631 f	Salmone 3	T TGTA C A

		42 (+)	ente rica serovar Typhi	AACC T CIGCIG G TIGG A GACGAC AT T IGICAC C
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CTACCTGC u	hp NC_003143 f rom 45227 90 to 45233 80 (-)	Yersini ³ a pes tis	T - TGCC C A CC A TGCTGCTG G T GG A ACGACGAC AT T T
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CTAGCCCT R	rom 46395 01 to 46413 78 (-)	onas aerugin osa P A01	C AGCC TGCTGCT A G TTGG ACGACGA AT T TGTCAC C
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CTCCTGCT s GCTG	NC_002947 f sb rom 5710 27 to 57157 2 (+)	Pseudom 3 onas putida KT244 0	T C CC A TGCTGCTG G GG ACGACGAC AT TTT TGTCAC
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CTCGGGTT f TCTTGCTG L	na NC_002929 f rom 30858 65 to 30984 55 (+)	Bordete 3 lla p ertussi s	T G GTT CT C C A G T TGCTGCTG G G A T A ACGACGAC AT TTT G GTC C-
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CTGACCTT p	NC_002947 f sta rom 8916 25 to 89371 2 (-)	Pseudom 3 onas putida KT244 0	T T C A C GACC TGCTGCT G TTGG ACGACGA AT T TGTCAC C
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CTGCCTGC o	NC_003143 f ern rom 3783 31 to 37887 6 (+)	Yersini ³ a pes tis	T C A C G CC TGCTGCT A G T GG ACGACGA AT T T TGTCAC C T - G T A
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	TGCTT D	63 to 21227 4 (+)	a sui s 1330	C G C TGCTGCT " G T G ACGACGA AT T T GTGTCAC C
GAM35 Hum CAGCAGCA 3678 an CACIGIGG TITGIA	CTTACTIG S CTGCTC B	NC_004431 f rom 29224 56 to 29232 41 (-)	Escheri 3 chia coli CFT073	IT C A C ACT TGCTGCT A G TGG ACGACGA AT TT TGTCAC C
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CIICCIGC d	NC_000922 f def rom 12217 35 to 12222	pneumon iae	T C CC TGCTGCT A G GG ACGACGA AT ITI TGTCAC C
GAM35 Hum CAGCAGCA 3678 an CACTGTGG TTTGTA	CTTCCTGC d	NC_002491 f Hef rom 12180 69 to 12186 29 (+)	Chlamyd 3 ophil a pneumon iae J138	C CC TGCTGCT A G GG ACGACGA AT TIT TGTCAC C

Replace paragraph 0159 with the following paragraph.

It is appreciated that the specific functions and accordingly the utilities of each of a plurality of GAM oligonucleotides that are described by Fig. 1 are correlated with and may be deduced from the

identity of the GAM TARGET GENES inhibited thereby, and whose functions are set forth in Table 8, hereby incorporated herein. Table 8, lines 685695-687709 shows data relating to the function and utilities of GAM RNA as set forth in SEO ID NO: 348.

After paragraph 0159, add the following Table 8.

GAM NAME	GAM RNA SEQUENCE	GAM ORGAN ISM	TAR	TARGET ORGANISM	GAM FUNCTION	GAM
	CACCAGCA CACTGTGG TITGTA	Human	ac cC	Bordetell a pertuss is	GAM53678 is a human mIRNA-like oligonuclectide, which torgets biotin carboxylase (accc., No.02429 from \$26407 carboxylase (accc., No.02429 from \$26407 carboxylase (accc., No.02429 from \$26407 carboxylase april of an anti-bacterial host defense mecha nism. accC BINDING SITE 1 and accC BINDING SITE 2 are bacterial taret binding sites that are found in the untranslated regions of sRNA encoded by the accC gene, corresponding to target binding sites such as BINDING SITE 1. BINDING SITE 1. BINDING SITE 1 and acc BINDING SITE 1. C. BINDING SITE 1. T. BINDING SITE 2. T. BINDING SITE 3. T. BINDING SITE 4. T. BINDING SITE 5. T. BINDING SITE 5	è
	CAGGAGCA CACTGTGG TTTGTA	Human	ac ex	Paeudomon as putida KT2440	GAM535678 is a human miRNA-like oligonucleotide, which targets isocitrate dehydrogensæe kinase/phosphatase (acek, NC_002947 from 5186742 to 5186457 (-1)), a backerial target gene enco ded by part of an ant i-backerial heat defense mechanism, acek BINDINS SITE 1 and acek BINDINS SITE 2 are bacterial hær defense mechanism. acek BINDINS SITE 2 are bacterial target binding sites that are found in the untranslated regions of miRNA encoded by the acek gene, corresponding to target binding sites that are found in the site of the site o	3

GAM353678 include t he diagnosis, prevention and treatment of Pseudomonas putida K T2440 infection and associated clinical conditions

Human acoR Pseudomon GAM35 CAGCAGCA 3678 CACTGTGG as aerugi nosa PA01

GAM353678 is a human miRNA-like oligonucleotide, which targets transcriptional regulator AcoR (acoR. NC_002516 from 463950 1 to 4641378)), a bacterial target gene encoded by the Ps eudomonas aeruginosa PA01 genome, as part of an anti-bacterial host defense mechanism, acoR BINDING SITE 1 and acoR BINDING SITE 2 are bacterial targ et binding sites that are found in the untranslated regions of mRNA encoded by the acoR gene, corresponding to target bindin q sites such as BINDING SITE I, BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of acoR BINDING SITE 1 and acoR BINDING SITE 2, and the complementary secondary st ructure to the nucleotide sequence of GAM353678 RNA are set fo rth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit acoR, a GAM353678 bacterial target gene which is associated with Pseudomonas aer uginosa PA01 infection, as part of an anti-bacterial host defe nse mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Pseudomonas aerugi nosa PA01 infection and associated clinical conditions

GAM35 CAGCAGCA Human aer-2 Pseudomon GAM353678 is a human miRNA-like 3678 CACTGTGG as putida KT2440

receptor Aer-2 (aer-2, NC_002947 from 2406996 to 2408561 (-)), a bacterial target gene encoded by the Pseudom onas putida KT2440 genome, as part of an antibacterial host d efense mechanism. aer-2 BINDING SITE 1 and aer-2 BINDING SITE 2 are bacterial ta rget binding sites that are found in the untranslated regions of mRNA encoded by the aer-2 gene, corresponding to target bin ding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of aer-2 BINDING SITE 1 and aer-2 BINDING SITE 2, and the complementary seconda ry structure to the nucleotide sequence of GAM353678 RNA are s et forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit aer-2, a GAM353678 bacterial target gene which is associated with Pseudomonas pu tida KT2440 infection, as part of an anti-bacterial host defen se mechanism. Accordingly, the utilities of GAM353678 include the diagnosis. prevention and treatment of Pseudomonas putida KT2440 infection and associated

oligonucleotide, which targets aerotaxis

Human amtB Shigella GAM35 CAGCAGCA 3678 CACTGTGG flexneri 2a str. 3

GAM353678 is a human miRNA-like oligonucleotide, which targets probable ammonium transporter (amtB, NC_004337 from 408059 to 409345 (+)), a bacterial target gene encoded by the Shi gella flexneri 2a str. 301 genome, as part of an anti-bacteria 1 host defense mechanism. amtB BINDING SITE 1 and amtB BINDING SITE

clinical conditions

2 are bacterial targ et binding sites that are found in the untranslated regions of mRNA encoded by the amtB gene. corresponding to target bindin g sites such as BINDING SITE I, BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of amtB BINDING SITE 1 and amtB BINDING SITE 2, and the complementary secondary at ructure to the nucleotide sequence of GAM353678 RNA are set fo rth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit amtB, a GAM353678 bacterial target gene which is associated with Shigella flexne ri 2a str. 301 infection, as part of an anti-bacterial host de fense mechanism. Accordingly, the utilities of GAM353678 inclu de the

diagnosis, prevention and treatment of Shigella flexmer i 2a str. 301 infection and associated clinical conditions

oligonucleotide, which targets Probable

GAM353678 is a human miRNA-like

GAM35 CAGCAGCA Human amtB Escherich 3678 CACTGTGG ia coli C

ammonium transporter (amtB, NC 004431 from 547616 to 548902 (+)), a bacterial target gene encoded by the Esc herichia coli CFT073 genome, as part of an antibacterial host defense mechanism, amtB BINDING SITE 1 and amtB BINDING SITE 2 are bacterial targ et binding sites that are found in the untranslated regions of mRNA encoded by the amtB gene, corresponding to target bindin g sites such as BINDING SITE I. BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of amtB BINDING SITE 1 and amtB BINDING SITE 2. and the complementary secondary st ructure to the nucleotide sequence of GAM353678 RNA are set fo rth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit amtB, a GAM353678 bacterial target gene which is associated with Escherichia col i CFT073 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Escherichia coli CFT07 3 infection and associated clinical conditions

GAM35 CAGCAGCA Human amtB Shigella 3678 CACTGTGG flexneri TTTGTA 2a str. 2 GAM353678 is a human miRNA-like oligonucleotide, which targets probable ammonium transporter (amtB, NC 004741 from 407860 to 409146 (+)), a bacterial target gene encoded by the Shi gella flexneri 2a str. 2457T genome, as part of an anti-bacter ial host defense mechanism. amtB BINDING SITE 1 and amtB BINDING SITE 2 are bacterial targ et binding sites that are found in the untranslated regions of mRNA encoded by the amtB gene. corresponding to target bindin g sites such as BINDING SITE I. BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of amtB BINDING SITE 1 and amtB BINDING SITE 2, and the complementary secondary st ructure to the nucleotide sequence of GAM353678 RNA are set fo rth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit amtB, a GAM353678

bacterial target gene which is associated with Shigella fizzer if 2m etc. 2m e

GAM35 CAGCAGCA Human aroA Neisseria 3678 CACTGTGG meningit TTTGTA idis Z249 GAM353678 is a human miRNA-like oligonucleotide, which targets 5enolpyruvoylshikimate-3-phosphate synthase (aroA, NC_003116 from 1557502 to (+)), a bacterial target gene enc oded by the Neisseria meningitidis Z2491 genome, as part of an antibacterial host defense mechanism. aroA BINDING SITE 1 and aroA BINDING SITE 2 are bacterial targ et binding sites that are found in the untranslated regions of mRNA encoded by the aroA gene, corresponding to target bindin q sites such as BINDING SITE I, BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of aroA BINDING SITE 1 and aroA BINDING SITE 2, and the complementary secondary st ructure to the nucleotide sequence of GAM353678 RNA are set fo rth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit aroA, a GAM353678 bacterial target gene which is associated with Neisseria menin gitidis Z2491 infection, as part of an anti-bacterial host def ense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Neisseria meningi tidis Z2491 infection and associated clinical conditions

GAM35 CAGCAGCA Human 3678 CACTGTGG TTTGTA

Human aroD Leptospir a interro gans sero var lai s tr. 56601

GAM353678 is a human miRNA-like oligonucleotide, which targets 3dehydroquinate dehydratase (aroD, NC_004342 from 48128 to 48832)), a bacterial target gene encoded by the Lept ospira interrogans serovar lai str. 56601 genome, as part of a n antibacterial host defense mechanism. aroD BINDING SITE 1 and aroD BINDING SITE 2 are bacterial targ et binding sites that are found in the untranslated regions of mRNA encoded by the aroD gene, corresponding to target bindin g sites such as BINDING SITE I, BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of aroD BINDING SITE 1 and aroD BINDING SITE 2. and the complementary secondary st ructure to the nucleotide sequence of GAM353678 RNA are set fo rth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit aroD, a GAM353678 bacterial target gene which is associated with Leptospira inte rrogans serovar lai str. 56601 infection, as part of an anti-b acterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Leptospira interrogans serovar lai str. 56601 infection and as sociated clinical conditions

GAM35 CAGCAGCA Human aroH Shigella 3678 CACTGTGG flexneri

Shigella GAM353678 is a human miRNA-like flexneri oligonucleotide, which targets 3-deoxy-D-2a str. 3 arabinoheptulosonate-7-phosphate synthase

(DAHP syn thetase, tryptophan repressible) (aroH, NC_004337 from 15575 27 to 1558573 (-)), a bacterial target gene encoded by the S higella flexneri 2a str. 301 genome, as part of an anti-bacterial host defense mechanism. aroH BINDING SITE 1 and aroH BINDING SITE 2 are bacterial targ et binding sites that are found in the untranslated regions of mRNA encoded by the aroH gene, corresponding to target bindin g sites such as BINDING SITE I. BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of aroH BINDING SITE 1 and aroH BINDING SITE 2, and the complementary secondary st ructure to the nucleotide sequence of GAM353678 RNA are set fo rth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit aroH, a GAM353678 bacterial target gene which is associated with Shigella flexne ri 2a str. 301 infection, as part of an anti-bacterial host de fense mechanism. Accordingly, the utilities of GAM353678 inclu de the diagnosis, prevention and treatment of Shigella flexner i 2a str. 301 infection and associated clinical conditions

GAM35 CAGCAGCA Human avtA Shigella 3678 CACTGTGG TTTGTA

flexneri 2a str. 3

GAM353678 is a human miRNA-like oligonucleotide, which targets alaninealpha-ketoisovalerate (or valine-pyruvate) transamina se, transaminase C (avtA, NC 004337 from 3721175 to 3722533

(+)), a bacterial target gene encoded by the Shigella flexne ri 2a str. 301 genome, as part of an anti-bacterial host defen se mechanism, avtA BINDING SITE is a bacterial target binding site that is a found in the the 3' untranslated region of mRNA encoded by th e avtA gene, corresponding to a target binding site such as BI NDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. T he nucleotide sequences of avtA BINDING SITE, and the compleme ntary secondary structure to the nucleotide sequence of GAM353 678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit avtA, a GAM353678 bacterial target gene which is associated with Shigella flexne ri 2a str. 301 infection, as part of an anti-bacterial host de fense mechanism. Accordingly, the utilities of GAM353678 inclu de the diagnosis, prevention and treatment of Shigella flexner i 2a str. 301 infection and associated clinical conditions

GAM35 CAGCAGCA 3678 CACTGTGG Human avtA Shigella flexneri 2a str. 2 457T

GAM353678 is a human miRNA-like oligonucleotide, which targets alaninealpha-ketoisovalerate/valine-pyruvate transaminase C (avtA, NC_004741 from

4052685 to 4053938 (-)), a bacterial target gene encoded by the Shigella flexneri 2a str. 2457T ge nome, as part of an anti-bacterial host defense mechanism. avtA BINDING SITE is a bacterial target binding site that is a found in the the 3' untranslated region of mRNA encoded by th e avtA gene, corresponding to a target binding site such as BI NDING SITE I, BINDING SITE II or BINDING SITE III of

Fig. 1. T he nucleotide sequences of avtA BINDING SITE, and the compleme ntary secondary structure to the nucleotide sequence of GAM353 678 RNA are set forth in Tables 6-7, hereby incorporated herei n. Another function of GAM353678 is to inhibit avtA, a GAM353678 bacterial target gene which is associated with Shigella flexne ri 2a str. 2457T infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 inc lude the diagnosis. prevention and treatment of Shigella flexn eri 2a str. 2457T infection and associated clinical conditions

GAM35 CAGCAGCA Human cII Salmonell 3678 CACTGTGG TTTGTA

a enteric a enteric a serovar Typhi Ty

GAM353678 is a human miRNA-like oligonucleotide, which targets transcriptional regulatory protein (cII, NC_004631 from 452 0121 to 4520342

(+)), a bacterial target gene encoded by the Salmonella enterica enterica serovar Typhi Ty2 genome, as par t of an antibacterial host defense mechanism. cII BINDING SITE 1 and CII BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of m RNA encoded by the cII gene, corresponding to target binding s ites such as BINDING SITE I, BINDING SITE II or BINDING SITE I II of Fig. 1. The nucleotide seguences of cII BINDING SITE 1 a nd cII BINDING SITE 2, and the complementary secondary structu re to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit cII, a GAM353678 b acterial target gene which is associated with Salmonella enterica enterica serovar Typhi Ty2 infection, as part of an anti-b acterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Salmonella enterica enterica serovar Typhi Ty2 infection and a ssociated clinical conditions

GAM35 CAGCAGCA 3678 CACTGTGG TTTGTA

a enteric a enteric a serovar Typhi

Human cII Salmonell GAM353678 is a human miRNA-like oligonucleotide, which targets transcriptional regulatory protein (cII, NC 003198 from 453 7312 to 4537533

(+)), a bacterial target gene encoded by the Salmonella enterica enterica serovar Typhi genome, as part of an antibacterial host defense mechanism, cII BINDING SITE 1 and cII BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of m RNA encoded by the cII gene, corresponding to target binding s ites such as BINDING SITE I. BINDING SITE II or BINDING SITE I II of Fig. 1. The nucleotide sequences of cII BINDING SITE 1 a nd cII BINDING SITE 2, and the complementary secondary structu re to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit cII, a GAM353678 b acterial target gene which is associated with Salmonella enter ica enterica serovar Typhi infection, as part of an anti-bacte rial host defense mechanism. Accordingly,

the utilities of GAM 353678 include the diagnosis, prevention and treatment of Salm onella enterica enterica serovar Typhi infection and associate d clinical

GAM35 CAGCAGCA Human cysM Salmonell 3678 CACTGTGG a typhimu rium LT2

GAM353678 is a human miRNA-like oligonucleotide, which targets cysteine synthase B (cysM, NC_003197 from 2551651 to 255256 2 (-)), a bacterial target gene encoded by the Salmonella ty phimurium LT2 genome, as part of an antibacterial host defens e mechanism. cysM BINDING SITE 1 and cysM BINDING SITE 2 are bacterial targ et binding sites that are found in the untranslated regions of mRNA encoded by the cysM gene, corresponding to target bindin g sites such as BINDING SITE I, BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of cysM BINDING SITE 1 and cysM BINDING SITE 2, and the complementary secondary st ructure to the nucleotide sequence of GAM353678 RNA are set fo rth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit cysM, a GAM353678 bacterial target gene which is associated with Salmonella typh imurium LT2 infection, as part of an anti-bacterial host defen se mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Salmonella typhimur ium LT2 infection and associated clinical conditions

GAM35 CAGCAGCA 3678 CACTGTGG KT2440 TTTGTA

Human cvsO Pseudomon GAM353678 is a human miRNA-like as putida oligonucleotide, which targets 3'(2'),5'bisphosphate nucleotidase (cysQ, NC_002947 from 315184 to 315984 (+)), a bacterial target gene encoded by t he Pseudomonas putida KT2440 genome, as part of an antibacter ial host defense mechanism. cysQ BINDING SITE 1 and cysQ BINDING SITE 2 are bacterial targ et binding sites that are found in the untranslated regions of mRNA encoded by the cysQ gene, corresponding to target bindin g sites such as BINDING SITE I. BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of cysQ BINDING SITE 1 and cysQ BINDING SITE 2, and the complementary secondary st ructure to the nucleotide sequence of GAM353678 RNA are set fo rth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit cysQ, a GAM353678 bacterial target gene which is associated with Pseudomonas put ida KT2440 infection. as part of an anti-bacterial host defens e mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Pseudomonas putida K T2440 infection and associated

GAM35 CAGCAGCA Human dadX Escherich 3678 CACTGTGG ia coli C TTTGTA

clinical conditions GAM353678 is a human miRNA-like oligonucleotide, which targets Alanine racemase, catabolic (dadX, NC_004431 from 1476306 to 1477376 (+)), a bacterial target gene encoded by the Esche richia coli CFT073 genome, as part of an antibacterial host d efense mechanism. dadX BINDING SITE 1 and dadX BINDING SITE 2 are bacterial targ et binding sites that are

found in the untranslated regions of mRNA encoded by the dadX gene, corresponding to target bindin g sites such as BINDING SITE I. BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide seguences of dadX BINDING SITE 1 and dadX BINDING SITE 2. and the complementary secondary st ructure to the nucleotide sequence of GAM353678 RNA are set fo rth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit dadX, a GAM353678 bacterial target gene which is associated with Escherichia col i CFT073 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Escherichia coli CFT07 3 infection and associated

GAM35 CAGCAGCA 3678 CACTGTGG TTTGTA

Human def

Chlamydop hila pneu moniae CW LO29

neu oli CW Po

GAM353678 is a human miRNA-like oligonucleotide, which targets Polypeptide Deformylase (def, NC_000922 from 1221735 to 122 2295 (+)), a bacterial target gene encoded by the Chlamydoph ila pneumoniae CWL029 genome, as part of an anti-bacterial host defense mechanism, def BINDING SITE 1 and def BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the def gene, corresponding to target binding sites such as BINDING SITE I. BINDING SITE II or BINDING SITE I II of Fig. 1. The nucleotide sequences of def BINDING SITE 1 and def BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit def, a GAM353678 b acterial target gene which is associated with Chlamydophila pn eumoniae CWL029 infection, as part of an anti-bacterial host d efense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Chlamydophila p neumoniae CWL029 infection and associated clinical conditions.

GAM35 CAGCAGCA 3678 CACTGTGG TTTGTA Human def Chlamydop hila pneu moniae J1

GAM353678 is a human miRNA-like oligonucleotide, which targets polypeptide deformylase (def, NC 002491 from 1218069 to 121 8629 (+)), a bacterial target gene encoded by the Chlamydoph ila pneumoniae J138 genome.as part of an anti-bacterial host defense mechanism, def BINDING SITE 1 and def BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of m RNA encoded by the def gene, corresponding to target binding s ites such as BINDING SITE I, BINDING SITE II or BINDING SITE I II of Fig. 1. The nucleotide sequences of def BINDING SITE 1 a nd def BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit def, a GAM353678 b acterial target gene which is associated

with Chlamydophile pn eumoniae J138 infection, as part of an anti-bacterial host def ense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Chlamydophila pne umoniae J138 infection and associated clinical conditions

GAM35 CAGCAGCA Human dnaE Mycobacte 3678 CACTGTGG rium lepr TTTGTA ae GAM353678 is a human miRNA-like oligonucleotide, which targets DNA polymerase III, [alpha] subunit (dnaE, NC_002677 from 1423014 to 1426547 (+)), a bacterial target gene encoded by the Mycobacterium leprae genome, as part of an anti-bacterial host defense mechanism. dnaE BINDING SITE 1 through dnaE BINDING SITE 3 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the dnaE gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of dnaE BINDING SITE 1 through dnaE BINDING SITE 3, and the complementary seco ndary structure to the nucleotide sequence of GAM353678 RNA ar e set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit dnaE, a GAM353678 bacterial target gene which is associated with Mycobacterium 1 eprae infection, as part of an anti-bacterial host defense med hanism, Accordingly, the utilities of GAM353678 include the diagnosis. prevention and treatment of Mycobacterium leprae infection and associated clinical conditions

GAM35 CAGCAGCA Human dsdA Salmonell 3678 CACTGTGG a typhimu TTTGTA rium LT2 GAM353678 is a human miRNA-like oligonucleotide, which targets D-serine deaminase (dsdA, NC_003197 from 4004453 to 4005775 (+)), a bacterial target gene encoded by the Salmonella typ himurium LT2 genome, as part of an anti-bacterial host defense mechanism. dsdA BINDING SITE 1 and dsdA BINDING SITE 2 are bacterial targ et binding sites that are found in the untranslated regions of mRNA encoded by the dsdA gene, corresponding to target bindin q sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of dsdA BINDING SITE 1 and dsdA BINDING SITE 2. and the complementary secondary st ructure to the nucleotide sequence of GAM353678 RNA are set fo rth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit dsdA, a GAM353678 bacterial target gene which is associated with Salmonella typh imurium LT2 infection, as part of an anti-bacterial host defen se mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Salmonella typhimurium LT2 infection and associated clinical conditions

GAM35 CAGCAGCA Human fepC Pseudomon 3678 CACTGTGG as aerugi TTTGTA nosa PA01 ASSOCIATED THIND A CONTINUE AND ASSOCIATED A CONTINUE A CONTINUE A CONTINUE AND ASSOCIATED ASSOCIAT

part of an an ti-bacterial host defense mechanism, fepC BINDING SITE 1 and fepC BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the fepC gene, corresponding to target bindin g sites such as BINDING SITE I. BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of fepC BINDING SITE 1 and fepC BINDING SITE 2. and the complementary secondary st ructure to the nucleotide sequence of GAM353678 RNA are set fo rth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit fepC, a GAM353678 bacterial target gene which is associated with Pseudomonas aer uginosa PA01 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Pseudomonas aerugi nosa PA01 infection and associated clinical conditions

GAM35 CAGCAGCA Human fhaL Bordetell GAM353678 is a human miRNA-like 3678 CACTGTGG a pertuss oligonucleotide, which targets TTGTA is (fhaL, NC 002929 from 3085865

a pertuss oligonucleotide, which targets adhesin is (fhaL, NC_002929 from 3085865 to 3098455 (+)), a bacterial target gene encoded by the Bordetella pertussis genome, as part of an anti-bacterial host defense mechanism, fhat BINDING SITE 1 and fhat BINDING SITE 2 are bacterial targ et binding sites that are found in the untranslated regions of mRNA encoded by the fhal gene, corresponding to target bindin g sites such as BINDING SITE I. BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of fhal-BINDING SITE 1 and fhal BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit fhal, a GAM353678 bacterial target gene which is associated with Bordetella pert ussis infection, as part of an anti-bacterial host defense med hanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Bordetella pertussis infection and associated clinical conditions

GAM35 CAGCAGCA Human flhB Bordetell
3678 CACTGTGG a pertuss
TTTGTA is

GAM353678 is a human miRNA-like oligonucleotide, which targets flagellar biosynthetic protein FlhB (flhB, NC 002929 from 1 441767 to 1442921 (+)), a bacterial target gene encoded by t he Bordetella pertussis genome, as part of an anti-bacterial h ost defense mechanism. flhB BINDING SITE 1 through flhB BINDING SITE 3 are bacterial target binding sites that are found in the untranslated region s of mRNA encoded by the flhB gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of flhB BINDING SITE 1 through flhB BINDING SITE 3, and the complementary seco ndary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby

incorporated herein. Another function of GAM353678 is to inhibit IRB, a GAM353678 bacterial target gene which is associated with Bordetella pert usels infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Bordetella pertussis infection and associated clinical conditions

GAM35 CAGCAGCA 3678 CACTGTGG Human fmtB(Staphyloc mr p) occus aur eus subsp

N315

GAM353678 is a human miRNA-like oligonucleotide, which targets FmtB protein (fmtB(mrp), NC_002745 from

2218145 to 2225590 (-)), a bacterial target gene encoded by the Staphylococcus aureus subsp. aureus N315 genome, as part of an anti-bacterial host defense mechanism. fmtB(mrp) BINDING SITE 1 and fmtB(mrp) BINDING SITE 2 are bact erial target binding sites that are found in the untranslated regions of mRNA encoded by the fmtB(mrp) gene, corresponding t o target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of fm tB(mrp) BINDING SITE 1 and fmtB(mrp) BINDING SITE 2. and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporate d herein. Another function of GAM353678 is to inhibit fmtB(mrp), a GAM35 3678 bacterial target gene which is associated with Staphyloco ccus aureus subsp. aureus N315 infection, as part of an anti-b acterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Staphylococcus aureus subsp. aureus N315 infection and associated clinical conditions

GAM35 CAGCAGCA 3678 CACTGTGG TTTGTA Human fmtB(Staphyloc mr p) occus aur eus subsp . aureus Mu50

GAM353678 is a human miRNA-like oligonucleotide, which targets FmtB protein (fmtB(mrp), NC_002758 from 2287935 to 2295380 (-)), a bacterial target gene encoded by the Staphylococcus aureus subsp. aureus Mu50 genome, as part of an anti-bacterial host defense mechanism. fmtB(mrp) BINDING SITE 1 and fmtB(mrp) BINDING SITE 2 are bact erial target binding sites that are found in the untranslated regions of mRNA encoded by the fmtB(mrp) gene, corresponding to target binding sites such as BINDING SITE I. BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of fm tB(mrp) BINDING SITE 1 and fmtB(mrp) BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporate d herein. Another function of GAM353678 is to inhibit fmtB(mrp), a GAM35 3678 bacterial target gene which is associated with Staphylococcus aureus subsp. aureus Mu50 infection, as part of an anti-b acterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Staphylococcus aureus subsp.

aureus Mu50 infection and associated clinical conditions

GAM35 CAGCACCA Human ftsY Chlamydop GAM353678 is a human miRNA-like 3678 CACTGTGG hila pneu oligonucleotide, which targets TTGTA moniae Jl division protein ftsY (ftsY, NC,

oligonucleotide, which targets cell division protein ftsY (ftsY, NC 002491 from 1113127 to 1113999 (-)), a bacterial target gene encoded by the Chlamy dophila pneumoniae J138 genome, as part of an anti-bacterial host defense mechanism. ftsY BINDING SITE 1 and ftsY BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the ftsY gene, corresponding to target bindin g sites such as BINDING SITE I, BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of ftsY BINDING SITE 1 and ftsY BINDING SITE 2, and the complementary secondary st ructure to the nucleotide sequence of GAM353678 RNA are set fo rth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit ftsY, a GAM353678 bacterial target gene which is associated with Chlamydophila pneumoniae J138 infection, as part of an anti-bacterial host de fense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Chlamydophila pn eumoniae J138 infection and associated clinical conditions

GAM35 CAGCAGCA Human ftsY Chlamydop 3678 CACTGTGG hila pneu TTTGTA moniae CW GAM353678 is a human miRNA-like oligonucleotide, which targets Cell Division Protein FtsY (ftsY, NC_000922 from 1115427 to 1116299 (-)), a bacterial target gene encoded by the Chlamy dophila pneumoniae CWL029 genome. as part of an anti-bacterial host defense mechanism. ftsY BINDING SITE 1 and ftsY BINDING SITE 2 are bacterial targ et binding sites that are found in the untranslated regions of mRNA encoded by the ftsY gene, corresponding to target bindin q sites such as BINDING SITE I. BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of ftsY BINDING SITE 1 and ftsY BINDING SITE 2, and the complementary secondary st ructure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit ftsY, a GAM353678 bacterial target gene which is associated with Chlamydophila pneumoniae CWL029 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Chlamydophila pneumoniae CWL029 infection and associated clinical conditions GAM353678 is a human miRNA-like

GAM35 CAGCAGCA Human gad Pseudomon 3678 CACTGTGG as putida TTTGTA KT2440

aminohydrolasme (gad, NC_002947 from 4871625 to 4872 929 (+)), a batterial target gene encoded by the Presudomonas putida KT2440 genome, an part of an antibacterial host defense mechanism. gad BINDINS SIZE 1 and gad BINDINS SIZE 2 are bacterial target binding sites that are found in the untranslated regions of m RWA

oligonucleotide, which targets guanine

encoded by the gad gene, corresponding to target binding sites such as BINDING SITE I. BINDING SITE II or BINDING SITE I II of Fig. 1. The nucleotide seguences of gad BINDING SITE 1 and gad BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth i n Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit gad, a GAM353678 b acterial target gene which is associated with Pseudomonas puti da KT2440 infection. as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Pseudomonas putida KT 2440 infection and associated clinical conditions

GAM35 CAGCAGCA 3678 CACTGTGG TTTGTA

Human glcC Escherich ia coli C

GAM353678 is a human miRNA-like oligonucleotide, which targets Glc operon transcriptional activator (glcC, NC 004431 from 3542871 to 3543695 (+)), a bacterial target gene encoded by the Escherichia coli CFT073 genome, as part of an anti-bacteri al host defense mechanism. glcC BINDING SITE 1 and glcC BINDING SITE 2 are bacterial targ et binding sites that are found in the untranslated regions of mRNA encoded by the glcC gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of glcC BINDING SITE 1 and glcC BINDING SITE 2, and the complementary secondary st ructure to the nucleotide sequence of GAM353678 RNA are set fo rth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit glcC. a GAM353678 bacterial target gene which is associated with Escherichia coli CFT073 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Escherichia coli CFT07 3 infection and associated clinical conditions

GAM35 CAGCAGCA 3678 CACTGTGG TTTGTA

Human qlqP Salmonell GAM353678 is a human miRNA-like a enteric oligonucleotide, which targets glycogen a enteric phosphorylase (glgP, NC 004631 from a serovar Typhi Ty

4129215 to 413 1662 (+)), a bacterial target gene encoded by the Salmonella enterica enterica serovar Typhi Ty2 genome, as part of an anti-bacterial host defense mechanism, glgP BINDING SITE 1 and qlqP BINDING SITE 2 are bacterial targ et binding sites that are found in the untranslated regions of mRNA encoded by the glgP gene, corresponding to target bindin g sites such as BINDING SITE I. BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of glgP BINDING SITE 1 and glgP BINDING SITE 2, and the complementary secondary st ructure to the nucleotide sequence of GAM353678 RNA are set fo rth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit glgP, a GAM353678

bacterial target gene which is associated with Salmonella ente rica enterica serovar Typhi Ty2 infection, as part of an antibacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Salmonella enterica enterica serovar Typhi Ty2 infection and associated clinical conditions

GAM35 CAGCAGCA Human glgP Salmonell 3678 CACTGTGG a enteric a enteric a serovar

Typhi

pestis KI

GAM353678 is a human miRNA-like oligonucleotide, which targets glycogen phosphorylase (glgP, NC_003198 from

4144568 to 414 7015 (+)), a bacterial target gene encoded by the Salmonella enterica enterica serovar Typhi genome, as part of an anti-bacterial host defense mechanism. glgP BINDING SITE 1 and glgP BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the glgP gene, corresponding to target bindin q sites such as BINDING SITE I, BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of qlqP BINDING SITE 1 and qlqP BINDING SITE 2, and the complementary secondary st ructure to the nucleotide sequence of GAM353678 RNA are set fo rth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit glgP, a GAM353678 bacterial target gene which is associated with Salmonella enterica enterica serovar Typhi infection, as part of an anti-bact erial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Sal monella enterica enterica serovar Typhi infection and associat ed clinical conditions

GAM35 CAGCAGCA Human glpC Yersinia 3678 CACTGTGG

GAM353678 is a human miRNA-like oligonucleotide, which targets snglycerol-3-phosphate dehydrogenase (anaerobic), K-small su bunit (qlpC, NC_004088 from 454677 to 456047 (+)), a bac terial target gene encoded by the Yersinia pestis KIM genome, as part of an anti-bacterial host defense mechanism. glpC BINDING SITE 1 and glpC BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the glpC gene, corresponding to target bindin g sites such as BINDING SITE I, BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of glpC BINDING SITE 1 and glpC BINDING SITE 2, and the complementary secondary st ructure to the

nucleotide sequence of GAM353678 RNA are set fo rth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit glpC, a GAM353678 bacterial target gene which is associated with Yersinia pestis KIM infection, as part of an anti-bacterial host defense mech anism. Accordingly, the utilities of GAM353678 include the dia gnosis, prevention and treatment of Yersinia pestis KIM infect ion and associated clinical conditions

GAM35 CAGCAGCA Human qlpC Yersinia 3678 CACTGTGG pestis

GAM353678 is a human miRNA-like oligonucleotide, which targets anaerobic glycerol-3-phosphate dehydrogenase subunit C (glpC, NC_003143 from 4289650 to 4290897 (-1), a bacterial target gene encoded by the Yersinia pestis genome, as part of an an ti-bacterial host defense mechanism, glpC BINDING SITE 1 and glpC BINDING SITE 2 are bacterial targ et binding sites that are found in the untranslated regions of mRNA encoded by the glpC gene, corresponding to target bindin g sites such as BINDING SITE I. BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of qlpC BINDING SITE 1 and glpC BINDING SITE 2, and the complementary secondary st ructure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit glpC, a GAM353678 bacterial target gene which is associated with Yersinia pestis infection, as part of an anti-bacterial host defense mechanis m. Accordingly, the utilities of GAM353678 include the diagnos is, prevention and treatment of Yersinia pestis infection and associated clinical conditions

GAM35 CAGCAGCA Human glpD Brucella 3678 CACTGTGG suis 1330

GAM353678 is a human miRNA-like oligonucleotide, which targets glycerol-3-phosphate dehydrogenase (glpD, NC_004310 2 10763 to 212274 (+)), a bacterial target gene encoded by th e Brucella suis 1330 genome, as part of an anti-bacterial host defense mechanism. glpD BINDING SITE 1 and glpD BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the glpD gene. corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of glpD BINDING SITE 1 and glpD BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set fo rth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit qlpD, a GAM353678 bacterial target gene which is associated with Brucella suis 1 330 infection, as part of an anti-bacterial host defense mecha nism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Brucella suis 1330 infection and associated clinical

GAM35 CAGCAGCA Human ipaH_ Shigella 3678 CACTGTGG 5 flexneri TTTGTA 2a str. 2 457T GAM353678 is a human miRNA-like oligonucleotide, which targets invasion plasmid antigen (ipaH 5, NC 004741 from 2023205 to 2024848 (+)), a bacterial target gene encoded by the Shigel la flexneri 2a str. 2457T genome, as part of an anti-bacterial host defense mechanism. ipaH 5 BINDING SITE 1 and ipaH 5 BINDING SITE 2 are bacterial target binding sites that are found in the untranslated region s of mRNA encoded by the ipaH_5 gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDI NG SITE III of Fig. 1. The nucleotide sequences of ipaH 5 BIND ING SITE 1 and ipaH 5 BINDING SITE 2, and the

complementary se condary structure to the nucleotide sequence of GAM353678 NAS near set forth in Tables 6-7, hereby incorporated herein. Another function of GAM53678 is to inhibit ipal. 5, a GAM53678 is to inhibit ipal. 5, a GAM53678 is to inhibit ipal. 5, a GAM53678 is not seen that the condition of GAM53678 is to inhibit ipal. 5, a GAM53678 in the condition of the condition o

GAM35 CAGCAGCA Hum 3678 CACTGTGG

Human ipaH9 Shigella .8 flexneri 2a str. 3 GAM353678 is a human miRNA-like oligonucleotide, which targets invasion plasmid antigen (ipaH9.8, NC_004337 from 1422064 to 1423779 (-)), a bacterial

target gene encoded by the Shige lla flexneri 2a str. 301 genome, as part of an anti-bacterial host defense mechanism. ipaH9.8 BINDING SITE 1 and ipaH9.8 BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regi ons of mRNA encoded by the ipaH9.8 gene, corresponding to target binding sites such as BINDING SITE I. BINDING SITE II or BI NDING SITE III of Fig. 1. The nucleotide sequences of ipaH9.8 BINDING SITE 1 and ipaH9.8 BINDING SITE 2, and the complementa ry secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit ipaH9.8, a GAM3536 78 bacterial target gene which is associated with Shigella fle xneri 2a str. 301 infection, as part of an antibacterial host defense mechanism. Accordingly, the utilities of GAM353678 in clude the diagnosis, prevention and treatment of Shigella flex neri 2a str. 301 infection and associated clinical

GAM35 CAGCAGCA 3678 CACTGTGG Human livH Bordetell a pertuss conditions. GAM353678 is a human miRNA-like oligonucleotide, which targets highaffinity branched-chain amino acid transport system perm ease protein (livH, NC 002929 from 1144729 to 1145607 , a bacterial target gene encoded by the Bordetella pertussis genome, as part of an anti-bacterial host defense mechanism. livH BINDING SITE is a bacterial target binding site that is a found in the the 3' untranslated region of mRNA encoded by th e livH gene, corresponding to a target binding site such as BI NDING SITE I. BINDING SITE II or BINDING SITE III of Fig. 1. T he nucleotide sequences of livH BINDING SITE, and the compleme ntary secondary structure to the nucleotide sequence of GAM353 678 RNA are set forth in Tables 6-7, hereby incorporated herei n. Another function of GAM353678 is to inhibit livH, a GAM353678 bacterial target gene which is associated with Bordetella pert ussis infection, as part of an antibacterial host defense mec hanism. Accordingly, the utilities of GAM353678 include the di agnosis, prevention and

GAMSS CAGCAGCA Human 1ppI Mycobacte 3678 CACTGTGG rium tube TTTGTA rculosis

treatment of Bordetella pertussis infe ction and associated clinical conditions

GAM353678 is a human miRNA-like

oligonucleotide, which targets lppI (lppI, NC 000962 from 2291267 to 2291923 (+)), a bac terial target gene encoded by the Mycobacterium tuberculosis H 37Rv genome, as part of an anti-bacterial host defense mechani sm. lppI BINDING SITE 1 and lppI BINDING SITE 2 are bacterial targ et binding sites that are found in the untranslated regions of mRNA encoded by the lppI gene, corresponding to target bindin q sites such as BINDING SITE I, BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of lppI BINDING SITE 1 and 1ppl BINDING SITE 2, and the complementary secondary st ructure to the nucleotide sequence of GAM353678 RNA are set fo rth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit lppI, a GAM353678 bacterial target gene which is associated with Mycobacterium t uberculosis H37Rv infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 in clude the diagnosis, prevention and treatment of Mycobacterium tuberculosis H37Rv infection and associated clinical conditio

GAM35 CAGCAGCA Human lppI Mycobacte 3678 CACTGTGG TTTGTA

rium bovi s subsp b ovis AF21 22/97

GAM353678 is a human miRNA-like oligonucleotide, which targets Probable

lipoprotein lppI (lppI, NC_002945 from 2275182 to 2275838 (+)), a bacterial target gene encoded by the Mycobac terium bovis subsp bovis AF2122/97 genome, as part of an anti- bacterial host defense mechanism. lppI BINDING SITE 1 and lppI BINDING SITE 2 are bacterial targ et binding sites that are found in the untranslated regions of mRNA encoded by the lppI gene, corresponding to target bindin q sites such as BINDING SITE I. BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of lppI BINDING SITE 1 and lppI BINDING SITE 2, and the complementary secondary st ructure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit lppI, a GAM353678 bacterial target gene which is associated with Mycobacterium bovis subsp bovis AF2122/97 infection, as part of an antibacte rial host defense mechanism. Accordingly, the utilities of GAM 353678 include the diagnosis, prevention and treatment of Myco bacterium bovis subsp bovis AF2122/97 infection and associated clinical conditions

GAM35 CAGCAGCA Human MGAT5 Human 3678 CACTGTGG

GAM353678 is a human miRNA-like oligonucleotide, which targets a human mannosyl (alpha-1,6-)-glycoprotein beta-1,6-N-acetyl- glucosaminyltransferase (MGAT5, Accession number: NM_002410) a s part of a host response mechanism associated with a Salmonel la typhimurium LT2 infection. MGAT5 BINDING SITE is a human target binding site that is a fo und in the the 3' untranslated region of mRNA encoded by the M GAT5 gene, corresponding to a target binding site such as BIND ING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. Add itionally, using the binding site prediction system of the pre sent invention GAM353678-A binds to sequences on orthologous UTR of rat(NM 023095). The nucleotide sequences of MGAT5 BINDIN G SITE, and the complementary secondary structure to the nucle otide sequence of GAM353678 RNA are set forth in Tables 6-7, h ereby incorporated herein. Another function of GAM353678 is to inhibit MGAT5, a GAM353678 human target gene which encodes an enzyme that catalyzes beta 1-6 branching on Nlinked carbohydrates. MGAT5 is associated with Salmonella typhimurium LT2 infection, and therefore GAM35 3678 is associated with the abovementioned infection, as part of a host response mechanism. Accordingly, the utilities of GA M353678 include the diagnosis, prevention and treatment of Sal monella typhimurium LT2 infection and associated clinical cond itions. The function of MGAT5 and its association with various diseases and clinical conditions has been established by previous stu dies. as described hereinabove with reference to GAM3451

GAM35 CAGCAGCA 3678 CACTGTGG TTTGTA tis

Human miaA Chlamydia GAM353678 is a human miRNA-like trachoma oligonucleotide, which targets tRNA isopentenylpyrophosphate transferase (miaA, NC_000117 from 899276 to 900295

(+)), a bacterial target gene encode d by the Chlamydia trachomatis genome, as part of an anti-bact erial host defense mechanism. miaA BINDING SITE 1 and miaA BINDING SITE 2 are bacterial targ et binding sites that are found in the untranslated regions of mRNA encoded by the miaA gene, corresponding to target bindin q sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of miaA BINDING SITE 1 and miaA BINDING SITE 2, and the complementary secondary st ructure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit miaA, a GAM353678 bacterial target gene which is associated with Chlamydia trach omatis infection, as part of an anti-bacterial host defense me chanism. Accordingly, the utilities of GAM353678 include the d iagnosis. prevention and treatment of Chlamydia trachomatis in fection and associated clinical conditions

GAM35 CAGCAGCA Human minE Pseudomon 3678 CACTGTGG as putida KT2440 TTTGTA

GAM353678 is a human miRNA-like oligonucleotide, which targets cell division topological specificity factor MinE (minE, NC_0 02947 from 1932680 to 1932934 (-)), a bacterial target gen e encoded by the Pseudomonas putida KT2440 genome, as part of an anti-bacterial host defense mechanism. minE BINDING SITE is a bacterial target binding site that is a found in the the 3' untranslated region

of mRNA encoded by the minE gene, corresponding to a target binding site such as BI NDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. T he nucleotide sequences of minE BINDING SITE. and the complementary secondary structure to the nucleotide sequence of GAM353 678 RNA are set forth in Tables 6-7, hereby incorporated herei n. Another function of GAM353678 is to inhibit minE, a GAM353678 bacterial target gene which is associated with Pseudomonas put ida KT2440 infection, as part of an anti-bacterial host defens e mechanism. Accordingly, the utilities of GAM353678 include t he diagnosis, prevention and treatment of Pseudomonas putida K T2440 infection and associated clinical conditions

GAM35 CAGCAGCA 3678 CACTGTGG TTTGTA Human nicT Mycobacte rium tube rculosis

GAM353678 is a human miRNA-like oligonucleotide, which targets nicT (nicT, NC_000962 from 3166681 to 3167799 (+)), a bac terial target gene encoded by the Mycobacterium tuberculosis H 37Rv genome, as part of an anti-bacterial host defense mechani sm. nicT BINDING SITE 1 and nicT BINDING SITE 2 are bacterial targ et binding sites that are found in the untranslated regions of mRNA encoded by the nicT gene, corresponding to target bindin g sites such as BINDING SITE I. BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of nicT BINDING SITE 1 and nicT BINDING SITE 2, and the complementary secondary st ructure to the nucleotide sequence of GAM353678 RNA are set fo rth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit nicT, a GAM353678 bacterial target gene which is associated with Mycobacterium t uberculosis H37Rv infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Mycobacterium tuberculosis H37Rv infection and associated clinical conditio

GAM35 CAGCAGCA 3678 CACTGTGG TTTGTA

Human nicT Mycobacte rium bovi s subsp b ovis AF21

GAM353678 is a human miRNA-like oligonucleotide, which targets POSSIBLE NICKEL-TRANSPORT INTEGRAL MEMBRANE PROTEIN NICT (nic T, NC_002945 from 3123200 to 3124318 (+)), a bacterial tar get gene encoded by the Mycobacterium boyis subsp bovis AF2122 /97 genome, as part of an anti-bacterial host defense mechanis m. nicT BINDING SITE 1 and nicT BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the nicT gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of nicT BINDING SITE 1 and nicT BINDING SITE 2, and the complementary secondary st ructure to the nucleotide sequence of GAM353678 RNA are set fo rth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit nicT, a GAM353678 bacterial target gene which is associated

with Mycobacterium b ovis subsp bovis AF2122/97 infection, as part of an antibacte rial host defense mechanism. Accordingly, the utilities of GAM 353678 include the diagnosis, prevention and treatment of Myco bacterium bovis subsp bovis AF2122/97 infection and associated clinical conditions

GAM35 CAGCAGCA 3678 CACTGTGG TTTGTA Human nupC Shigella flexneri 2a str. 3 GAM353678 is a human miRNA-like oligonucleotide, which targets permease of transport system for 3 nucleosides (nupC, NC_0043 37 from 2515842 to 2517083 (+)), a bacterial target gene encoded by the Shigella flexneri 2a str. 301 genome. as part of an anti-bacterial host defense mechanism, nupC BINDING SITE 1 through nupC BINDING SITE 3 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the nupC gene, corresponding to target bi nding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of nupC BINDING SITE 1 through nupC BINDING SITE 3, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA ar e set forth in Tables 6-7, hereby incorporated herein, Another function of GAM353678 is to inhibit nupC. a GAM353678 bacterial target gene which is associated with Shigella flexne ri 2a str. 301 infection, as part of an antibacterial host defense mechanism. Accordingly, the utilities of GAM353678 inclu de the diagnosis, prevention and treatment of Shigella flexner i 2a str. 301 infection and associated clinical conditions

GAM35 CAGCAGCA 3678 CACTGTGG TTTGTA Human nupC Escherich ia coli C GAM353678 is a human miRNA-like oligonucleotide, which targets Nucleoside permease nupC (nupC, NC_004431 from 2795390 to 2 796631 (+)), a bacterial target gene encoded by the Escheric hia coli CFT073 genome, as part of an antibacterial host defe nse mechanism. nupC BINDING SITE 1 through nupC BINDING SITE 3 are bacterial target binding sites that are found in the untranslated region s of mRNA encoded by the nupC gene, corresponding to target bi nding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of nupC BINDING SITE 1 through nupC BINDING SITE 3, and the complementary seco ndary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit nupC, a GAM353678 bacterial target gene which is associated with Escherichia coli CFT073 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Escherichia coli CFT07 3 infection and associated clinical conditions

GAM35 CAGCAGCA 3678 CACTGTGG Human nupC Shigella flexneri 2a str. 2

Shigella GAM353678 is a human miRNA-like flexneri oligonucleotide, which targets permease 2a str. 2 of transport system for 3 nucleosides 457T

(nupC, NC 0047 41 from 2494019 to 2495221 (+)), a bacterial target gene e ncoded by the Shigella flexneri 2a str. 2457T genome, as part of an anti-bacterial host defense mechanism, nupC BINDING SITE 1 through nupC BINDING SITE 3 are bacterial target binding sites that are found in the untranslated region s of mRNA encoded by the nupC gene, corresponding to target bi nding sites such as BINDING SITE I. BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of nupC BINDING SITE 1 through nupC BINDING SITE 3, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit nupC, a GAM353678 bacterial target gene which is associated with Shigella flexne ri 2a str. 2457T infection, as part of an antibacterial host defense mechanism. Accordingly, the utilities of GAM353678 inc lude the diagnosis, prevention and treatment of Shigella flexn eri 2a str. 2457T infection and associated clinical conditions

GAM35 CAGCAGCA Human ompG Escherich 3678 CACTGTGG ia coli C TTTGTA FT073

GAM353678 is a human miRNA-like oligonucleotide, which targets Outer membrane protein G precursor (ompG, NC 004431 from 16 24577 to 1625533

(+)), a bacterial target gene encoded by th e Escherichia coli CFT073 genome, as part of an anti-bacterial host defense mechanism. ompG BINDING SITE 1 and ompG BINDING SITE 2 are bacterial targ et binding sites that are found in the untranslated regions of mRNA encoded by the ompG gene, corresponding to target bindin g sites such as BINDING SITE I, BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of ompG BINDING SITE 1 and ompG BINDING SITE 2, and the complementary secondary st ructure to the nucleotide sequence of GAM353678 RNA are set fo rth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit ompG, a GAM353678 bacterial target gene which is associated with Escherichia coli CFT073 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis. prevention and treatment of Escherichia coli CFT07 3 infection and associated clinical conditions

GAM35 CAGCAGCA Human orn Yersinia 3678 CACTGTGG pestis TTTGTA GAMSIS678 is a human miRNA-like oligonucleotide, which targets oligoribonuclease (orn, NC_003149 from J9831 to J98876 (+)), a bacterial target gene encoded by the Yersinia pestis genome, as part of an anti-bacterial host defense mechanism. orn BINDING SITE is a bacterial target binding site that is a found in the the 3' untranslated region of miNA encoded by the orn gene, corresponding to a target binding site such as BIND ING SITE I; BINDING SITE II or BINDING SIT

and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-T, hereby incorporated herein. Another function of GAM353678 is to inhibit orn, a GAM353678 and acterial target open which is associated with Yersinia pestis infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM354678 conclude the diagnosis, prevention and treatment of Yersinia pestis infection and associated quintage initial conditions.

GAM35 CAGCAGCA 3678 CACTGTGG TTTGTA Human oxyR Salmonell a enteric a enteric

Typhi

GAM353678 is a human miRNA-like oligonucleotide, which targets hydrogen peroxide-inducible regulon activator (oxyR, NC_00319 8 from 3607204 to 3608121 (-)), a bacterial target gene en coded by the Salmonella enterica enterica serovar Typhi genome , as part of an antibacterial host defense mechanism. oxyR BINDING SITE 1 and oxyR BINDING SITE 2 are bacterial targ et binding sites that are found in the untranslated regions of mRNA encoded by the oxyR gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of oxyR BINDING SITE 1 and oxyR BINDING SITE 2. and the complementary secondary st ructure to the nucleotide sequence of GAM353678 RNA are set fo rth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit oxyR, a GAM353678 bacterial target gene which is associated with Salmonella enterica enterica serovar Typhi infection, as part of an anti-bact erial host defense mechanism. Accordingly, the utilities of GA M353678 include the diagnosis, prevention and treatment of Sal monella enterica enterica serovar Typhi infection and associat ed clinical

GAM35 CAGCAGCA 3678 CACTGTGG TTTGTA Human oxyR Salmonell a typhimu rium LT2 conditions GAM353678 is a human miRNA-like oligonucleotide, which targets oxidative stress regulatory protein (oxyR, NC_003197 from 4343080 to 4343997 (+)), a bacterial target gene encoded by the Salmonella typhimurium LT2 genome, as part of an anti-bacte rial host defense mechanism. oxvR BINDING SITE 1 and oxvR BINDING SITE 2 are bacterial targ et binding sites that are found in the untranslated regions of mRNA encoded by the oxyR gene. corresponding to target bindin g sites such as BINDING SITE I, BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of oxvR BINDING SITE 1 and oxyR BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set fo rth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit oxyR, a GAM353678 bacterial target gene which is associated with Salmonella typh imurium LT2 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Salmonella typhimur ium LT2 infection and

GAM35 CAGCAGCA 3678 CACTGTGG TTTGTA o gerewar Typhi Ty

associated clinical conditions Human oxvR Salmonell GAM353678 is a human miRNA-like a enteric oligonucleotide, which targets hydrogen a enteric peroxide-inducible regulon activator (oxvR, NC 00463 1 from 3592864 to 3593781 (-)), a bacterial target gene en coded by the Salmonella enterica enterica

serovar Typhi Ty2 genome, as part of an anti-bacterial host defense mechanism. oxyR BINDING SITE 1 and oxyR BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the oxyR gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of oxyR BINDING SITE 1 and oxyR BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set fo rth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit oxyR, a GAM353678

bacterial target gene which is associated with Salmonella ente rica enterica serovar Typhi Ty2 infection, as part of an antibacterial host defense mechanism. Accordingly, the utilities o f GAM353678 include the diagnosis, prevention and treatment of Salmonella enterica enterica serovar Typhi Ty2 infection and associated clinical conditions

GAM35 CAGCAGCA 3678 CACTGTGG TTTGTA KT2440

Human pbpG Pseudomon GAM353678 is a human miRNA-like as putida oligonucleotide, which targets D-alanyl-D-alanine-endopeptidase (pbpG, NC_002947 from 4323 707 to 4324633 (+)), a bacterial target gene encoded by the Pseudomonas putida KT2440 genome, as part of an anti-bacterial host defense mechanism. pbpG BINDING SITE 1 and pbpG BINDING SITE 2 are bacterial targ et binding sites that are found in the untranslated regions of mRNA encoded by the pbpG gene, corresponding to target bindin q sites such as BINDING SITE I, BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of pbpG BINDING SITE 1 and pbpG BINDING SITE 2, and the complementary secondary st ructure to the nucleotide sequence of GAM353678 RNA are set fo rth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit pbpG, a GAM353678 bacterial target gene which is associated with Pseudomonas put ida KT2440 infection. as part of an anti-bacterial host defens e mechanism. Accordingly, the utilities of GAM353678 include t he diagnosis, prevention and treatment of Pseudomonas putida K T2440 infection and associated

GAM35 CAGCAGCA Human pchA Pseudomon 3678 CACTGTGG as aerugi TTTGTA nosa PA01

clinical conditions GAM353678 is a human miRNA-like oligonucleotide, which targets salicylate biosynthesis isochorismate synthase (pchA, NC_0025 16 from 4745120 to 4746550

(+)), a bacterial target gene e ncoded by the Pseudomonas aeruginosa PA01 genome, as part of a n anti-bacterial host defense mechanism. pchA BINDING SITE 1 and pchA BINDING SITE 2 are bacterial targ et

binding sites that are found in the untranslated regions of mRNA encoded by the pchA gene, corresponding to target bindin g sites such as BINDING SITE I. BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide seguences of pchA BINDING SITE 1 and pchA BINDING SITE 2. and the complementary secondary st ructure to the nucleotide sequence of GAM353678 RNA are set fo rth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit pchA, a GAM353678 bacterial target gene which is associated with Pseudomonas aer uginosa PA01 infection, as part of an anti-bacterial host defe nse mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Pseudomonas aerugi nosa PA01 infection and associated clinical conditions

GAM35 CAGCAGCA Human pcnA Mycobacte GAM353678 is a human miRNA-like 3678 CACTGTGG rium lepr oligonucleotide, which targets ae (pcnA, NC, 002677 from 324825)

oligonucleotide, which targets pcnA (pcnA, NC_002677 from 3248268 to 3249728 (-)), a bac terial target gene encoded by the Mycobacterium leprae genome, as part of an anti-bacterial host defense mechanism, pcnA BINDING SITE 1 and pcnA BINDING SITE 2 are bacterial targ et binding sites that are found in the untranslated regions of mRNA encoded by the pcnA gene, corresponding to target bindin g sites such as BINDING SITE I, BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of pcnA BINDING SITE 1 and ponA BINDING SITE 2. and the complementary secondary st ructure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit pcnA, a GAM353678 bacterial target gene which is associated with Mycobacterium 1 eprae infection, as part of an anti-bacterial host defense med hanism. Accordingly, the utilities of GAM353678 include the di agnosis, prevention and treatment of Mycobacterium leprae infection and associated clinical conditions

GAM35 CACCAGCA Human phnV Salmonell GAM353678 is a human minNA-like
3678 CACTOTG a tenter o diponucleotide, which targets
1TIGTA a enteric enterior component of 2a server aminoethiphosphomate transport

Typhi

oligonucleotide, which targets probable membrane component of 2aminoethylphosphonate transp orter (phnV. NC 003198 from 471575 to 472366)), a bac terial target gene encoded by the Salmonella enterica enterica serovar Typhi genome, as part of an anti-bacterial host defen se mechanism, phnV BINDING SITE 1 and phnV BINDING SITE 2 are bacterial targ et binding sites that are found in the untranslated regions of mRNA encoded by the phnV gene, corresponding to target bindin g sites such as BINDING SITE I, BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of phnV BINDING SITE 1 and phnV BINDING SITE 2, and the complementary secondary st ructure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit phnV, a GAM353678

bacterial target gene which is associated with Salmonella ente rica enterica serovar Typhi infection, as part of an anti-bact erial host defense mechanism, Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Sal monella enterica enterica serovar Typhi infection and associated clinical conditions

GAM35 CAGCAGCA 3678 CACTGTGG

a enteric a serovar Typhi Ty

Human phnV Salmonell GAM353678 is a human miRNA-like oligonucleotide, which targets probable a enteric membrane component of 2-

aminoethylphosphonate transp orter (phnV, NC 004631 from 2508735 to 2509526 (+)), a bac terial target gene encoded by the Salmonella enterica enterica serovar Typhi Ty2 genome, as part of an antibacterial host defense mechanism. phnV BINDING SITE 1 and phnV BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the phnV gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of phnV BINDING SITE 1 and phnV BINDING SITE 2, and the complementary secondary st ructure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit phnV, a GAM353678 bacterial target gene which is associated with Salmonella ente rica enterica serovar Typhi Ty2 infection, as part of an antibacterial host defense mechanism. Accordingly, the utilities o f GAM353678 include the diagnosis, prevention and treatment of Salmonella enterica enterica serovar Typhi Ty2 infection and associated clinical conditions

GAM35 CAGCAGCA Human phoY2 Mycobacte 3678 CACTGTGG rium bovi

s subsp b ovis AF21 GAM353678 is a human miRNA-like oligonucleotide, which targets PROBABLE PHOSPHATE-TRANSPORT SYSTEM TRANSCRIPTIONAL REGULATOR Y PROTEIN PHOY2 (phoY2, NC_002945 from 914388 to 915029 (-)), a bacterial target gene encoded by the Mycobacterium bov is subsp bovis AF2122/97 genome, as part of an anti-bacterial host defense mechanism. phoY2 BINDING SITE 1 and phoY2 BINDING SITE 2 are bacterial ta rget binding sites that are found in the untranslated regions of mRNA encoded by the phoY2 gene, corresponding to target bin ding sites such as BINDING SITE I. BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of phoY2 BINDING SITE 1 and phoy2 BINDING SITE 2. and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are s et forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit phoY2, a GAM353678 bacterial target gene which is associated with Mycobacterium bovis subsp bovis AF2122/97 infection, as part of an antibact erial host defense mechanism. Accordingly, the utilities of GA M353678 include the diagnosis, prevention and treatment of Myc obacterium bovis subsp bovis AF2122/97 infection and associated

clinical conditions

GAM35 CAGCAGCA 3678 CACTGTGG TTTGTA

Human phoY2 Mycobacte GAM353678 is a human miRNA-like rium tube rculosis H37Rv

oligonucleotide, which targets phoY2 (phoY2, NC 000962 from 913556 to 914197 (-)), a b acterial target gene encoded by the Mycobacterium tuberculosis H37Rv genome, as part of an anti-bacterial host defense mecha nism. phoY2 BINDING SITE 1 and phoY2 BINDING SITE 2 are bacterial ta rget binding sites that are found in the untranslated regions of mRNA encoded by the phoY2 gene, corresponding to target bin ding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of phoY2 BINDING SITE 1 and phoY2 BINDING SITE 2, and the complementary seconda ry structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit phoY2, a GAM353678 bacterial target gene which is associated with Mycobacterium tuberculosis H37Rv infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Mycobacterium tuberculosis H37Rv infection and associated clinical conditions

GAM35 CAGCAGCA 3678 CACTOTOS TTTGTA

as putida KT2440

Human pill Pseudomon GAM353678 is a human miRNA-like oligonucleotide, which targets type IV pili twitching motility protein PilT (pilT, NC_002947 from 5816934 to 5817944 (-)), a bacterial target gene enc oded by the Pseudomonas putida KT2440 genome, as part of an an ti-bacterial host defense mechanism. pilT BINDING SITE 1 through pilT BINDING SITE 3 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the pilT gene, corresponding to target bi nding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of pilT BINDING SITE 1 through pilT BINDING SITE 3, and the complementary seco ndary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit pill. a GAM353678 bacterial target gene which is associated with Pseudomonas put ida KT2440 infection, as part of an anti-bacterial host defens e mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Pseudomonas putida K T2440 infection and associated clinical conditions

GAM35 CAGCAGCA 3678 CACTGTGG

Human polA Mycobacte rium lepr

GAM353678 is a human miRNA-like oligonucleotide, which targets DNA polymerase I (polA, NC_002677 from 1648220 to 1650955 (-)), a bacterial target gene encoded by the Mycobacterium le prae genome, as part of an anti-bacterial host defense mechani sm. polA BINDING SITE 1 and polA BINDING SITE 2 are bacterial targ et binding sites that are found in the untranslated regions of mRNA encoded by the polA gene, corresponding to target bindin q sites such as BINDING SITE I,

BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of polA BINDING SITE 1 and polA BINDING SITE 2. and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set fo rth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit pola, a GAM353678 bacterial target gene which is associated with Mycobacterium 1 eprae infection, as part of an anti-bacterial host defense med hanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Mycobacterium leprae infection and associated clinical conditions

GAM35 CAGCAGCA Human prcA Mycobacte 3678 CACTGTGG rium lepr TTTGTA ae GAM353678 is a human miRNA-like oligonucleotide, which targets proteasome [alpha]-type subunit 1 (prcA, NC_002677 1576553 to 1577350 (+)), a bacterial target gene encoded by the Mycobacterium leprae genome, as part of an anti-bacterial host defense mechanism. prcA BINDING SITE 1 and prcA BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the prcA gene. corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of proA BINDING SITE 1 and prcA BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit prcA, a GAM353678 bacterial target gene which is associated with Mycobacterium 1 eprae infection, as part of an anti-bacterial host defense mec hanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Mycobacterium leprae infection and associated clinical

GAM35 CAGCAGCA Human pta Pseudomon 3678 CACTGTGG as putida TTTGTA KT2440 conditions GAM353678 is a human miRNA-like oligonucleotide, which targets phosphate acetyltransferase (pta, NC_002947 from 891625 to 893712 (-)), a bacterial target gene encoded by the Pseudo monas putida KT2440 genome, as part of an antibacterial host defense mechanism. pta BINDING SITE 1 and pta BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the pta gene, corresponding to target binding sites such as BINDING SITE I. BINDING SITE II or BINDING SITE I II of Fig. 1. The nucleotide sequences of pta BINDING SITE 1 and pta BINDING SITE 2, and the complementary secondary structu re to the nucleotide sequence of GAM353678 RNA are set forth i n Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit pta, a GAM353678 b acterial target gene which is associated with Pseudomonas puti da KT2440 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis,

prevention and treatment of Pseudomonas putida KT 2440 infection and associated clinical conditions

GAM35 CAMCOACCA Human pteH Salmonell CAM353678 is a human miRNA-like
3678 CAMCOTOR

a enteric plagomuclectide, which targets
TITOTA a enteric phosphocarrier protein HPr [pt
a servers NC 003198 from 2505403 to 25

Typhi

oligonucleotide, which targets phosphocarrier protein HPr (ptsH, NC 003198 from 2505403 to 2505660 (+)), a bacterial target gene encoded by the Salmonella enterica enterica serovar Typhi genome, as part of an anti-bacterial host defense mechanism. ptsH BINDING SITE is a bacterial target binding site that is found in the the 3' untranslated region of mRNA encoded by the ptsH gene, corresponding to a target binding site such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. T he nucleotide sequences of ptsH BINDING SITE, and the compleme ntary secondary structure to the nucleotide sequence of GAM353 678 RNA are set forth in Tables 6-7, hereby incorporated herei n. Another function of GAM353678 is to inhibit ptsH, a GAM353678 bacterial target gene which is associated with Salmonella ente rica enterica serovar Typhi infection, as part of an anti-bact erial host defense mechanism, Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Sal monella enterica enterica serovar Typhi infection and associat ed clinical

GAM35 CAGCAGCA Human rbsR Shigella 3678 CACTETGS flexneri TTTGTA 2a str. 3 GAM353678 is a human miRNA-like oligonucleotide, which targets regulator for rbs operon (rbsR, NC_004337 from

3947708 to 3 948700 (+)), a bacterial target gene encoded by the Shigella flexneri 2a str. 301 genome, as part of an anti-bacterial host defense mechanism. rbsR BINDING SITE 1 through rbsR BINDING SITE 3 are bacterial target binding sites that are found in the untranslated region s of mRNA encoded by the rbsR gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of rbsR BINDING SITE 1 through rbsR BINDING SITE 3, and the complementary seco ndary structure to the nucleotide sequence of GAM353678 RNA ar e set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit rbsR, a GAM353678 bacterial target gene which is associated with Shigella flexne ri 2a str. 301 infection, as part of an anti-bacterial host de fense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Shigella flexner i 2a str. 301 infection and associated clinical conditions

GAM35 CAGCAGCA Human rbsR Shigella 3678 CACTGTGG flexneri TTTGTA 2a str. 2

oligonucleotide, which targets regulator for rbs operon (rbsR, Nc_004741 from 3824594 to 3 825577 (-)), a bacterial target gene encoded by the Shigella flexneri 2s str. 2457f genome, as part of an anti-bacterial h out defense mechanism. rbsR BINDING SITE 1 through rbsR BINDING SITE 3 rebacterial target binding sites

GAM353678 is a human miRNA-like

conditions

that are found in the untranslated region s of mRNA encoded by the rbsR gene, corresponding to target bi nding sites such as BINDING SITE I. BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of rbsR BINDING SITE 1 through rbsR BINDING SITE 3, and the complementary seco ndary structure to the nucleotide sequence of GAM353678 RNA ar e set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit rbsR, a GAM353678 bacterial target gene which is associated with Shigella flexne ri 2a str. 2457T infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Shigella flexn eri 2a str. 2457T infection and associated clinical conditions

GAM35 CAGCAGCA Human rbsR Escherich 3678 CACTGTGG ia coli C TTTGTA FT073

GAM353678 is a human miRNA-like oligonucleotide, which targets Ribose operon repressor (rbsR, NC 004431 from 4439260 to 44 40252 (+)), a bacterial target gene encoded by the Escherich ia coli CFT073 genome, as part of an antibacterial host defense mechanism, rbsR BINDING SITE 1 through rbsR BINDING SITE 3 are bacterial target binding sites that are found in the untranslated region s of mRNA encoded by the rbsR gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SITE III of Fig.1. The nucleotide sequences of rbsR BINDING SITE 1 through rbsR BINDING SITE 3, and the complementary seco ndary structure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit rbsR, a GAM353678 bacterial target gene which is associated with Escherichia col i CFT073 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Escherichia coli CFT07 3 infection and associated clinical conditions

GAM35 CAGCAGCA Human recG Mycobacte 3678 CACTGTGG rium lepr GAM353678 is a human miRNA-like oligonucleotide, which targets ATPdependent DNA helicase (recG, NC 002677 from 2014723 to 2016954 (-)), a bacterial target gene encoded by the Mycoba cterium leprae genome, as part of an anti-bacterial host defen se mechanism. recG BINDING SITE 1 and recG BINDING SITE 2 are bacterial targ et binding sites that are found in the untranslated regions of mRNA encoded by the reaG gene. corresponding to target bindin g sites such as BINDING SITE I, BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of recG BINDING SITE 1 and recG BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set fo rth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit recG, a GAM353678

bacterial target gene which is associated with Mycobacterium I eprae infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM535678 include the diagnosis, prevention and treatment of Mycobacterium leprae infection and associated clinical conditions.

GAM35 CAGCAGCA Human relA Mycobacte
3678 CACTGTGG rium bovi
TTTGTA subsp b
ovis AF21

22/97

GAM353678 is a human miRNA-like oligonucleotide, which targets PROBABLE GTP PYROPHOSPHOKINASE RELA (ATP:GTP 3'-PYROPHOSPHOTR ANSFERASE) (PPGPP SYNTHETASE ((P)PPGPP SYNTHETASE) (GTP DIP HOSPHOKINASE) (relA, NC_002945 from 2875274 to 2877646 (-)), a bacterial target gene encoded by the Mycobacterium bovis subsp bovis AF2122/97 genome, as part of an anti-bacterial host defense mechanism. relA BINDING SITE 1 and relA BINDING SITE 2 are bacterial targ et binding sites that are found in the untranslated regions of mRNA encoded by the relA gene, corresponding to target bindin q sites such as BINDING SITE I, BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of relA BINDING SITE 1 and relA BINDING SITE 2. and the complementary secondary st ructure to the nucleotide sequence of GAM353678 RNA are set fo rth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit relA, a GAM353678 bacterial target gene which is associated with Mycobacterium boyis subsp boyis AF2122/97 infection, as part of an antibacte rial host defense mechanism. Accordingly, the utilities of GAM 353678 include the diagnosis, prevention and treatment of Myco bacterium bovis subsp bovis AF2122/97 infection and associated clinical conditions

GAM35 CAGCAGCA Human relA Mycobacte 3678 CACTETGG rium tube TTTGTA roulosis

Human relA Mycobacte GAM353678 is a human miRNA-like rium tube oligonucleotide, which targets relA rculosis (relA, NC_000962 from 2907824 to 2910196

(-)), a bac terial target gene encoded by the Mycobacterium tuberculosis H 37Rv genome, as part of an anti-bacterial host defense mechani sm. relA BINDING SITE 1 and relA BINDING SITE 2 are bacterial targ et binding sites that are found in the untranslated regions of mRNA encoded by the relA gene, corresponding to target bindin g sites such as BINDING SITE I. BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of relA BINDING SITE 1 and relA BINDING SITE 2. and the complementary secondary st ructure to the nucleotide sequence of GAM353678 RNA are set fo rth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit relA, a GAM353678 bacterial target gene which is associated with Mycobacteriumt uberculosis H37Ry infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 in clude the diagnosis, prevention and treatment of Mycobacterium tuberculosis H37Rv infection and associated clinical conditio GAM35 CAGCAGCA 3678 CACTGTGG TTTGTA

Human risA Bordetell GAM353678 is a human miRNA-like a pertuss oligonucleotide, which targets tresponse regulator protein (risA, NC 002929 from target gene encoded by the Borde tella pertussis genome, as part of an antibacterial host defense mechanism, risA BINDING SITE 1 and risA BINDING SITE 2 are bacterial targ et binding sites that are found in the untranslated regions of mRNA encoded by the risk gene, corresponding to target bindin g sites such as BINDING SITE I, BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of risk BINDING SITE 1 and risk BINDING SITE 2, and the complementary secondary st ructure to the nucleotide sequence of GAM353678 RNA are set fo rth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit risA, a GAM353678 bacterial target gene which is associated with Bordetella pert ussis infection, as part of an anti-bacterial host defense med hanism. Accordingly, the utilities of GAM353678 include the diagnosis. prevention and treatment of Bordetella pertussis infection and associated clinical conditions

GAM35 CAGCAGCA 3678 CACTGTGG

Human rpsT Pseudomon GAM353678 is a human miRNA-like as putida KT2440

oligonucleotide, which targets ribosomal protein S20 (rpsT, NC_002947 from 707068 to 7073 46 (-)), a bacterial target gene encoded by the Pseudomonas putida KT2440 genome, as part of an anti-bacterial host defen se mechanism. rpsT BINDING SITE 1 and rpsT BINDING SITE 2 are bacterial targ et binding sites that are found in the untranslated regions of mRNA encoded by the rpsT gene, corresponding to target bindin g sites such as BINDING SITE I, BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of rpsT BINDING SITE 1 and rpsT BINDING SITE 2, and the complementary secondary st ructure to the nucleotide sequence of GAM353678 RNA are set fo rth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit rpsT, a GAM353678 bacterial target gene which is associated with Pseudomonas put ida KT2440 infection, as part of an anti-bacterial host defens e mechanism. Accordingly, the utilities of GAM353678 include the diagnosis. prevention and treatment of Pseudomonas putida K T2440 infection and associated clinical conditions

GAM35 CAGCAGCA Human ruvB Yersinia 3678 CACTGTGG pestis

GAM353678 is a human miRNA-like oligonucleotide, which targets Holliday junction DNA helicase (ruvB, NC_003143 from 233644 9 to 2337453 (+)), a bacterial target gene encoded by the Ye rsinia pestis genome, as part of an antibacterial host defens e mechanism. ruvB BINDING SITE 1 and ruvB BINDING SITE 2 are bacterial targ et binding sites that are found in the untranslated regions of mRNA encoded by the ruvB gene, corresponding to target bindin q sites such as BINDING SITE I, BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of ruvB

BINDING SITE 1 and ruvB BINDING SITE 2, and the complementary secondary st ructure to the nucleotide sequence of CAM555678 RNA are set for thin Tables 6-7, hereby incorporated herein. Another function of CAM55678 is to intinit ruva, a CAM55678 bacterial target gene which is associated with Yershin perils infection, as part of an anti-bacterial boat defended and the complete of th

GAM35 CAGCAGCA 3678 CACTGTGG Human ruvB Yersinia pestis KI GAM353678 is a human miRNA-like oligonucleotide, which targets Holliday junction helicase subunit A (ruyB. NC_004088 from 2482031 to 2483035 a bacterial target gene encoded by the Yersinia pestis KIM genome, as part of an anti-bacterial host defense mechanism. ruvB BINDING SITE 1 and ruvB BINDING SITE 2 are bacterial targ et binding sites that are found in the untranslated regions of mRNA encoded by the ruvB gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of ruvB BINDING SITE 1 and ruvB BINDING SITE 2, and the complementary secondary at ructure to the nucleotide sequence of GAM353678 RNA are set fo rth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit ruvB, a GAM353678 bacterial target gene which is associated with Yersinia pestis KIM infection, as part of an anti-bacterial host defense mech anism. Accordingly, the utilities of GAM353678 include the dia gnosis. prevention and treatment of Yersinia pestis KIM infection and associated

GAM35 CAGCAGCA 3678 CACTGTGG TTTGTA Human selB Pseudomon as putida KT2440

clinical conditions GAM353678 is a human miRNA-like oligonucleotide, which targets selenocysteine-specific translation elongation factor (selB, NC_002947 from 582133 to 584055 (+)), a bacterial target gene encoded by the Pseudomonas putida KT2440 genome, as part of an antibacterial host defense mechanism, selB BINDING SITE is a bacterial target binding site that is a found in the the 3' untranslated region of mRNA encoded by th e selB gene, corresponding to a target binding site such as BI NDING SITE I. BINDING SITE II or BINDING SITE III of Fig. 1. T he nucleotide seguences of selB BINDING SITE, and the compleme ntary secondary structure to the nucleotide sequence of GAM353 678 RNA are set forth in Tables 6-7, hereby incorporated herei n. Another function of GAM353678 is to inhibit selB, a GAM353678 bacterial target gene which is associated with Pseudomonas put ida KT2440 infection, as part of an anti-bacterial host defens e mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Pseudomonas putida K T2440 infection and associated clinical

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GAM35 CAGCAGCA Human SERPI Human 3678 CACTGTGG NH 1 TTTGTA

GAM353678 is a human miRNA-like oligonucleotide, which targets a human Serine proteinase inhibitor clade H (heat shock prote in 47) member 1; (SERPINH1, Accession number: NM 001235) as pa rt of a host response mechanism associated with a Escherichia coli CFT073, Streptococcus pneumoniae R6. Streptococcus pneumo niae TIGR4, Streptococcus pyogenes M1 GAS, Streptococcus pyoge nes MGAS315, Streptococcus pyogenes MGAS8232 and Streptococcus pyogenes SSI-1 infections. SERPINH1 BINDING SITE 1 and SERPINH1 BINDING SITE 2 are human target binding sites that are found in the untranslated region s of mRNA encoded by the SERPINH1 gene, corresponding to targe t binding sites such as BINDING SITE I, BINDING SITE II or BIN DING SITE III of Fig. 1. Additionally, using the binding site prediction system of the present invention GAM353678-A binds t o sequences on orthologous UTR of rat(NM 017173). The nucleoti de sequences of SERPINH1 BINDING SITE 1 and SERPINH1 BINDING S ITE 2, and the complementary secondary structure to the nucleo tide sequence of GAM353678 RNA are set forth in Tables 6-7, he reby incorporated herein. Another function of GAM353678 is to inhibit SERPINH1, a GAM353 678 human target gene which encodes a heat shock protein and s erpin, that may function as a chaperone for procollagen in the ER. SERPINH1 is associated with Escherichia coli CFT073. Stre ptococcus pneumoniae R6. Streptococcus pneumoniae TIGR4, Strep tococcus pyogenes M1 GAS, Streptococcus pyogenes MGAS315, Stre ptococcus pyogenes MGAS8232 and Streptococcus pyogenes SSI-1 i nfections, and therefore GAM353678 is associated with the abov ementioned infections, as part of a host response mechanism. A ccordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Escherichia coli CFT073, Streptoco ccus pneumoniae R6, Streptococcus pneumoniae TIGR4, Streptococ cus pyogenes M1 GAS, Streptococcus pyogenes MGAS315, Streptoco ccus pyogenes MGAS8232 and Streptococcus pyogenes SSI-1 infect ions and associated clinical conditions. The function of SERPINH1 and its association with various dise ases and clinical conditions has been established by previous studies, as described hereinabove with reference to

GAM35 CAGCAGCA Human sitD Shigella 3678 CACTGTGG flexneri TTTGTA 2a str. 3

GAM353678 is a human miRNA-like oligonucleotide, which targets Iron transport protein, inner membrane component (siCD, NC_00 4337 from 1 40836 to 140821 [9]). Short state of the size of

CAMRSO

bindin q sites such as BINDING SITE I, BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of sitD BINDING SITE 1 and sitD BINDING SITE 2. and the complementary secondary st ructure to the nucleotide sequence of GAM353678 RNA are set fo rth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit sitD, a GAM353678 bacterial target gene which is associated with Shigella flexne ri 2a str. 301 infection, as part of an anti-bacterial host de fense mechanism. Accordingly, the utilities of GAM353678 inclu de the diagnosis, prevention and treatment of Shigella flexner i 2a str. 301 infection and associated clinical conditions

GAM35 CAGCAGCA 3678 CACTGTGG Human sitD Shigella flexneri 2a str. 2 GAM353678 is a human miRNA-like oligonucleotide, which targets Iron transport protein, inner membrane component (sitD, NC_00 4741 from to 1905523 (+)), a bacterial target gene encoded by the Shigella flexneri 2a str. 2457T genome, as par t of an antibacterial host defense mechanism, sitD BINDING SITE 1 and sitD BINDING SITE 2 are bacterial targ et binding sites that are found in the untranslated regions of mRNA encoded by the sitD gene, corresponding to target binding sites such as BINDING SITE I. BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide seguences of sitD BINDING SITE 1 and sitD BINDING SITE 2. and the complementary secondary st ructure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit sitD, a GAM353678 bacterial target gene which is associated with Shigella flexne ri 2a str. 2457T infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 inc lude the diagnosis, prevention and treatment of Shigella flexn eri 2a str. 2457T infection and associated clinical conditions

GAM35 CAGCAGCA 3678 CACTGTGG TTTGTA Human speD Salmonell a enteric a enteric a serovar Typhi Ty

GAM353678 is a human miRNA-like oligonucleotide, which targets Sadenosylmethionine decarboxylase proenzyme (speD, NC_004631 from 196380 to 197174 (-)), a bacterial target gene encoded by the Salmonella enterica enterica serovar Typhi Ty2 gen ome, as part of an antibacterial host defense mechanism, speD BINDING SITE 1 and speD BINDING SITE 2 are bacterial targ et binding sites that are found in the untranslated regions of mRNA encoded by the speD gene, corresponding to target bindin g sites such as BINDING SITE I. BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of speD BINDING SITE 1 and speD BINDING SITE 2. and the complementary secondary st ructure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit speD, a GAM353678 bacterial target gene which is associated with Salmonella ente rica enterica serovar Typhi Ty2 infection, as part of an antibacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Salmonella enterica enterica serovar Typhi Ty2 infection and associated clinical conditions

GAM35 CAGCAGCA 3678 CACTGTGG Human speD Salmonell a enteric a enteric a serovar Typhi

GAM353678 is a human miRNA-like oligonucleotide, which targets Sadenosylmethionine decarboxylase proenzyme (speD, NC 003198 from 196389 to 197183 (-)), a bacterial target gene enc oded by the Salmonella enterica enterica serovar Typhi genome, as part of an antibacterial host defense mechanism. speD BINDING SITE 1 and speD BINDING SITE 2 are bacterial targ et binding sites that are found in the untranslated regions of mRNA encoded by the speD gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of speD BINDING SITE 1 and speD BINDING SITE 2, and the complementary secondary st ructure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit speD, a GAM353678 bacterial target gene which is associated with Salmonella enterica enterica serovar Typhi infection, as part of an anti-bact erial host defense mechanism. Accordingly. the utilities of GAM353678 include the diagnosis, prevention and treatment of Sal monella enterica enterica serovar Typhi infection and associated clinical conditions

GAM35 CAGCAGCA 3678 CACTGTGG Human speD Salmonell a typhimu rium LT2

GAM353678 is a human miRNA-like oligonucleotide, which targets Sadenosylmethionine decarboxylase (speD, NC_003197 from 1 94201 to 194995 (-)), a bacterial target gene encoded by th e Salmonella typhimurium LT2 genome, as part of an anti-bacter ial host defense mechanism. speD BINDING SITE 1 and speD BINDING SITE 2 are bacterial targ et binding sites that are found in the untranslated regions of mRNA encoded by the speD gene, corresponding to target binding sites such as BINDING SITE I. BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of speD BINDING SITE 1 and speD BINDING SITE 2. and the complementary secondary st ructure to the nucleotide sequence of GAM353678 RNA are set fo rth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit speD, a GAM353678 bacterial target gene which is associated with Salmonella typh imurium LT2 infection, as part of an anti-bacterial host defen se mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Salmonella typhimurium LT2 infection and associated clinical conditions

GAM35 CAGCAGCA 3678 CACTGTGG TTTGTA Human ssb Pseudomon as putida KT2440 GAM353678 is a human miRNA-like oligonucleotide, which targets single stranded DNA-binding protein (asb, NC_002947 from 5 71027 to 571572 (+)), a bacterial target gene encoded by th e Pseudomonas putida KT2440 genome, as part of an anti-bacteri al host defense mechanism, ssb BINDING SITE 1 and ssb BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of m RNA encoded by the ssb gene, corresponding to target binding s ites such as BINDING SITE I. BINDING SITE II or BINDING SITE I II of Fig. 1. The nucleotide sequences of ssb BINDING SITE 1 a nd ssb BINDING SITE 2, and the complementary secondary structu re to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit ssb, a GAM353678 b acterial target gene which is associated with Pseudomonas puti da KT2440 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Pseudomonas putida KT 2440 infection and associated

GAM35 CAGCAGCA 3678 CACTGTGG TTTGTA

Human sseB Escherich GAM353678 is a human miRNA-like oligonucleotide, which targets Protein sseB (sseB, NC 004431 from 2922456 to 2923241 (-)), a bacterial target gene encoded by the Escherichia coli CFT 073 genome, as part of an anti-bacterial host defense mechanism. sseB BINDING SITE 1 and sseB BINDING SITE 2 are bacterial targ et binding sites that are found in the untranslated regions of mRNA encoded by the sseB gene, corresponding to target bindin g sites such as BINDING SITE I. BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of sseB BINDING SITE 1 and sseB BINDING SITE 2. and the complementary secondary st ructure to the nucleotide sequence of GAM353678 RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit sseB, a GAM353678 bacterial target gene which is associated with Escherichia col i CFT073 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Escherichia coli CFT07 3 infection and associated

GAM35 CAGCAGCA 3678 CACTGTGG TTTGTA

a pertuss is

clinical conditions Human tcfA Bordetell GAM353678 is a human miRNA-like oligonucleotide, which targets tracheal colonization factor precursor (tcfA,

NC 002929 from 1264436 to 1266379

(+)), a bacterial target gene encoded b y the Bordetella pertussis genome, as part of an anti-bacteria 1 host defense mechanism, tcfA BINDING SITE 1 and tcfA BINDING SITE 2 are bacterial targ et binding sites that are found in the untranslated regions of mRNA encoded by the tcfA gene, corresponding to target bindin q sites such as BINDING SITE I, BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of tcfA BINDING SITE 1 and tofA BINDING SITE 2, and the complementary secondary st ructure to the nucleotide sequence of GAM353678

RNA are set forth in Tables 6-7, hereby incorporated herein. Another function of GAM533678 is to inhibit tota, a GAM533678 bacterial target gene which is associated with Bordetella pert ussis infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM533678 include the diagnosis, and the control of and treatment of Bordetella percentage of the control of the design of the design of the control of the design of the desig

GAM35 CAGCAGCA 3678 CACTGTGG TTTGTA

Human truA Mycobacte rium lepr

GAM353678 is a human miRNA-like oligonucleotide, which targets probable pseudouridylate synthase (truA, NC_002677 from 234 3329 to 2344078 (-)), a bacterial target gene encoded by the Mycobacterium leprae genome, as part of an anti-bacterial host defense mechanism. truA BINDING SITE 1 and truA BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the truA gene, corresponding to target bindin g sites such as BINDING SITE I, BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of truA BINDING SITE 1 and truA BINDING SITE 2, and the complementary secondary st ructure to the nucleotide sequence of GAM353678 RNA are set fo rth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit truA, a GAM353678 bacterial target gene which is associated with Mycobacterium 1 eprae infection, as part of an anti-bacterial host defense mec hanism. Accordingly, the utilities of GAM353678 include the di agnosis. prevention and treatment of Mycobacterium leprae infe ction and associated clinical conditions

GAM35 CAGCAGCA H 3678 CACTGTGG TTTGTA

Human trunc Staphyloc at ed occus aur fmtB eus subsp . aureus GAM353678 is a human miRNA-like oligonucleotide, which targets truncated FmtB protein (truncated fmtB, NC_003923 from 2238 083 to 2240143 (-)), a bacterial target gene encoded by the Staphylococcus aureus subsp. aureus MW2 genome, as part of an anti-bacterial host defense mechanism. truncated fmtB BINDING SITE 1 and truncated fmtB BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the truncated fmtB gene, corresponding to target binding sites such as BINDING SITE I. BINDING SITE II or BINDING SITE III of Fig. 1. The nucleotide sequences of truncated fmtB BINDING SITE 1 and truncated fmtB BINDING SITE 2, and the complementary secondary structure to t he nucleotide sequence of GAM353678 RNA are set forth in Table s 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit truncated fmtB, a GAM353678 bacterial target gene which is associated with Staph ylococcus aureus subsp. aureus MW2 infection, as part of an an ti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Staphylococcus aureus subsp. aureus MW2 infection and associated

GAM35 CAGCAGCA Human uhpA Yersinia 3678 CACTGTGG pestis TTTGTA

GAM353678 is a human miRNA-like oligonucleotide, which targets twocomponent system response regulator (uhpA. NC 003143 from 4522790 to 4523380 (-)). a bacterial target gene encoded by the Yersinia pestis genome, as part of an anti-bacterial host defense mechanism. uhpA BINDING SITE 1 and uhpA BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the uhpA gene, corresponding to target binding sites such as BINDING SITE I, BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of uhpA BINDING SITE 1 and uhpA BINDING SITE 2, and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set fo rth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit uhpA, a GAM353678 bacterial target gene which is associated with Yersinia pestis infection, as part of an anti-bacterial host defense mechanis m. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Yersinia pestis infection and

associated clinical conditions

GAM35 CAGCAGCA Human ung Haemophil GAM353678 is a human miRNA-like 3678 CACTGTGG ne influe nzae Rd

oligonucleotide, which targets uracil DNA glycosylase (ung, NC_000907 from to 1933 5 (+)), a bacterial target gene encoded by the Haemophilus influenzae Rd genome, as part of an anti-bacterial host defen se mechanism. ung BINDING SITE 1 and ung BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the ung gene, corresponding to target binding s ites such as BINDING SITE I, BINDING SITE II or BINDING SITE I II of Fig. 1. The nucleotide sequences of ung BINDING SITE 1 a nd ung BINDING SITE 2, and the complementary secondary structu re to the nucleotide sequence of GAM353678 RNA are set forth i n Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit ung, a GAM353678 b acterial target gene which is associated with Haemophilus infl uenzae Rd infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis. prevention and treatment of Haemophilus influenza e Rd infection and associated clinical conditions

GAM35 CAGCAGCA Human vanB Pseudomon GAM353678 is a human miRNA-like 3678 CACTGTGG as aerugi TTTGTA nosa PA01

oligonucleotide, which targets vanillate O-demethylase oxidoreductase (vanB, NC_002516 from 5504120 to 5505073

(+)), a bacterial target gene encoded b y the Pseudomonas aeruginosa PA01 genome, as part of an anti-b acterial host defense mechanism, vanB BINDING SITE 1 and vanB BINDING SITE 2 are bacterial target binding sites that are found in the untranslated regions of mRNA encoded by the vanB gene, corresponding to target bindin q sites such as BINDING SITE I,

BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of vanB BINDING SITE 1 and wanB BINDING SITE 2. and the complementary secondary structure to the nucleotide sequence of GAM353678 RNA are set fo rth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit vanB, a GAM353678 bacterial target gene which is associated with Pseudomonas aer uginosa PA01 infection, as part of an anti-bacterial host defe ase mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Pseudomonas aerugi nosa PA01 infection and associated clinical conditions

GAM35 CAGCAGCA Human yabO Escherich 3678 CACTGTGG ia coli C TTTGTA FT073 GAM353678 is a human miRNA-like oligonucleotide, which targets Ribosomal large subunit pseudouridine synthase A

(yab0, NC_00 4431 from 61489 to 62148 (-)), a bacterial target gene encoded by the Escherichia coli CFT073 genome, as part of an anti-bacterial host defense mechanism, yabo BINDING SITE 1 and yabo BINDING SITE 2 are bacterial targ et binding sites that are found in the untranslated regions of mRNA encoded by the yabO gene, corresponding to target bindin g sites such as BINDING SITE I. BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of yab0 BINDING SITE 1 and vabo BINDING SITE 2. and the complementary secondary st ructure to the nucleotide sequence of GAM353678 RNA are set fo rth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit yabo, a GAM353678 bacterial target gene which is associated with Escherichia col i CFT073 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis, prevention and treatment of Escherichia coli CFT07 3 infection and associated clinical conditions

GAM35 CAGCAGCA Human yciE Escherich 3678 CACTGTGG ia coli C TTTGTA FT073 GAM353678 is a human miRNA-like oligonucleotide, which targets Protein vciE (vciE, NC 004431 from 1558641 to 1559147 (-)), a bacterial target gene encoded by the Escherichia coli CFT 073 genome, as part of an anti-bacterial host defense mechanis m. vciE BINDING SITE 1 and vciE BINDING SITE 2 are bacterial targ et binding sites that are found in the untranslated regions of mRNA encoded by the yciE gene, corresponding to target bindin g sites such as BINDING SITE I. BINDING SITE II or BINDING SIT E III of Fig. 1. The nucleotide sequences of vciE BINDING SITE 1 and voie BINDING SITE 2. and the complementary secondary st ructure to the nucleotide sequence of GAM353678 RNA are set fo rth in Tables 6-7, hereby incorporated herein. Another function of GAM353678 is to inhibit yciE, a GAM353678 bacterial target gene which is associated with Escherichia col i CFT073 infection, as part of an anti-bacterial host defense mechanism. Accordingly, the utilities of GAM353678 include the diagnosis,

prevention and treatment of Escherichia coli CFT07 3 infection and associated clinical conditions

Replace paragraph 0160 with the following paragraph.

Studies documenting the well known correlations between each of a plurality of GAM TARGET GENEs that are described by Fig.1 and the known gene functions and related diseases are listed in Table 9, hereby incorporated herein. Specifically, in Table 9, lines 6046-6059 describes references of GAM target genes, as set forth in SEQ ID NO.348 in Table 8.

After paragraph 0160, add the following Table 9, paragraph, Table 11, paragraph, Table 12, paragraph, and Table 13.

Table 9:

TARGET	TARGET ORGANISM	REFERENCES
MGAT5	Human	Demetriou, M.; Granovsky, M.; Quaggin, S.; Dennis, J. W.: Negative regulation of T-cell activation and autoimmunity by Mgat5 N-glycosylation. Nature 409: 733-739, 2001.
MGAT5	Human	Granovsky, M.; Fata, J.; Pawling, J.; Muller, W. J.; Khokha, R.; Dennis, J. W.:Suppression of tumor growth and metastasis in Mgat5-deficient mice. Nature Med.6: 306-12, 2000.
MGAT5	Human	Saito, H.; Nishikawa, A.; Gu, J.; Ihara, Y.; Soejima, H.; Wada, Y.; Sekiya, C.; Niikawa, N.; Taniguchi, N.: cDNA cloning and chromosomal mapping of human N-acetyl glucosaminyltransferase V+. Biochem. Biophys. Res. Commun.

Table 11, lines 275482-275565, shows data of GAM RNA SEQ ID NO:348 printed on microarray chip probes, as described in detail in Fig.17.

Table 11

PROBE SEQUENCE	PROBE TYPE	GAM RNA S MIR NAME	EQ ID/ GAM	RNA/MIR	SEQUENCE		SIG Y NAL	BACKG ROUND Z-SCO RE	MISM ATCH Z-SCO RE
CCCAGCAGCAC ACTGTGGTTTG TACGCGATCCG TTATCGTTCGG TATCGAACGTA ACGAT	Predicted	348	CAGCAGCAC	ACTGTGGTT	TTGTA	A2	638	4.2	3.2
CCCAGCAGCAC	Predicted	348	CAGCAGCAC	ACTGTGGT1	TTGTA	D2	9435	16.6	20.9

ACTGTGGTTTG TACGCGATCCG TTATCGTTCGG TATCGAACGTA ACGAT							
CCCAGCAGCAC ACTGTGGTTTG TACGCGATCCG TTATCGTTCGG TATCGAACGTA ACGAT	Predicted	348	CAGCAGCACACTGTGGTTTGTA	F	25910	14.8	27.5
CCCAGCAGCAC ACTGTGGTTTG TACGCGATCCG TTATCGTTCGG TATCGAACGTA ACGAT	Predicted	348	CAGCAGCACACTGTGGTTTGTA	F1	65518	12.0	30.2
CCCAGCAGCAC ACTGTGGTTTG TACGCGATCCG TTATCGTTCGG TATCGAACGTA ACGAT	Predicted	348	CAGCAGCACACTGTGGTTTGTA	G1	65518	10.1	29.3
CCCAGCAGCAC ACTGTGGTTTG TACGCGATCCG TTATCGTTCGG TATCGAACGTA ACGAT	Predicted	348	CAGCAGCACACTGTGGTTTGTA	H1	37067	9.9	28.2
CCCAGCAGCAC ACTGTGGTTTG TACGGATCGTT ATAACGATCCG GTATCGAACGT AACGA	Predicted	348	CAGCAGCACACTGTGGTTTGTA	A2	606	3.7	3.2
CCCAGCAGCAC ACTGTGGTTTG TACGGATCGTT ATAACGATCCG GTATCGAACGT AACGA	Predicted	348	CAGCAGCACACTGTGGTTTGTA	D2	7549	15.4	19.5
CCCAGCAGCAC ACTGTGGTTTG TACGGATCGTT ATAACGATCCG GTATCGAACGT AACGA	Predicted	348	CAGCAGCACACTGTGGTTTGTA	E1	20239	13.8	25.3
CCCAGCAGCAC ACTGTGGTTTG TACGGATCGTT ATAACGATCCG GTATCGAACGT AACGA	Predicted	348	CAGCAGCACACTGTGGTTTGTA	F1	65518	12.0	29.3
CCCAGCAGCAC ACTGTGGTTTG TACGGATCGTT ATAACGATCGG GTATCGAACGT AACGA	Predicted	348	CAGCAGCACACTGTGGTTTGTA	G1	65518	10.1	28.0
CCCAGCAGCAC ACTGTGGTTTG	Predicted	348	CAGCAGCACACTGTGGTTTGTA	Н1	27597	9.2	25.8

TACGGATCGTT ATAACGATCCG GTATCGAACGT AACGA

Table 12, line 177, shows data relating to GAM RNA SEQ ID NO:348 that were validated by means of Wet Laboratory.

Table 12

GAM RNA SEQUENCE	VALIDATION METHOD	SIGNAL	BACKGROUND Z-SCORE	MISMATCH Z-SCORE	GAM RNA SEQ-ID
CAGCAGCACACTGTGGTTTGTA	Chip strong	65518	16.623587	30.172779	348

Table 13, lines 3-42, 47-69, 84-121, 143-179, 187-207, 210-256, 264-478 shows sequence data of GAMs associated with different bacterial infections.

Table 13

	NAME	SEQ ID NOS OF GAMS ASSOCIATED WITH INFECTION
2	Bordetella	$\begin{array}{c} 1, 6, \ 10, \ 11, \ 12, \ 13, \ 16, \ 18, \ 19, \ 20, \ 21, \ 22, \ 23, \ 24, \ 25, \ 26, \ 27, \ 28, \ 29, \ 33, \ 34, \ 37, \ 41, \ 42, \ 43, \ 44, \ 74, \ 84, \ 95, \ 52, \ 53, \ 54, \ 55, \ 75, \ 58, \ 59, \ 60, \ 63, \ 65, \ 66, \ 67, \ 68, \ 69, \ 70, \ 71, \ 75, \ 76, \ 77, \ 79, \ 84, \ 86, \ 87, \ 88, \ 89, \ 91, \ 94, \ 96, \ 97, \ 99, \ 90,$
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39 Yersinia

40 Yersinia pestis KIM